

GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: January 8, 2003, 16:40:45 / Search time 2182 Seconds

(without alignments)

11281.916 Million cell updates/sec

Title: US-09-847-665-4

Sequence: 1520

1 actcaaggaagaaatcattc.....cgtcagcgtccttgaana 1520

Scoring table:

IDENTITY_NUC
Gapop 10.0, Gapext 1.0

Number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database:

EST.*
1: em_estda.*
2: em_esthum.*
3: em_estin.*
4: em_estnu.*
5: em_estrov.*
6: em_estrpl.*
7: em_estro.*
8: em_hcc.*
9: gb_est1.*
10: gb_est2.*
11: gb_hcc.*
12: gb_hcc3.*
13: gb_est4.*
14: gb_est5.*
15: em_estfun.*
16: em_estom.*
17: gb_gss.*
18: em_gss_hum.*
19: em_gss_inv.*
20: em_gss_pln.*
21: em_gss_vit.*
22: em_gss_fun.*
23: em_gss_mam.*
24: em_gss_mus.*
25: em_gss_other.*
26: em_gss_pro.*
27: em_gss_rtd.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	378.2	24.9	391	9	A1694972
2	345.4	22.7	367	9	AA324132
3	327	21.5	474	9	AL120587
4	299.2	19.7	741	13	B1457956
5	299.2	19.7	785	12	BG114610
6	299.2	19.7	798	12	BG116320

7	261.6	17.2	586	13	B1602678	B1602678
8	164.4	10.8	459	14	BQ601726	BQ601726
9	82.2	5.4	1101	17	CNS00EVL	CNS00EVL
10	76.8	5.1	1101	17	CNS0039G	CNS0039G
11	75.4	5.0	905	17	CNS00KHX	CNS00KHX
12	74.2	4.9	1101	17	CNS00E07	CNS00E07
13	74	4.9	1101	17	CNS003B0	CNS003B0
14	73.6	4.8	975	17	BH179465	BH179465
15	72.8	4.8	1101	17	CNS004ZM	CNS004ZM
16	72.4	4.8	1167	17	CNS07360	CNS07360
17	71.8	4.7	524	17	CNS01U90	CNS01U90
18	71.6	4.7	987	17	CNS014PQ	CNS014PQ
19	71.4	4.6	1190	17	CNS020M7	CNS020M7
20	70.4	4.6	945	17	CNS04D0K	CNS04D0K
21	69.6	4.6	576	17	CNS035N7	CNS035N7
22	69.2	4.6	1001	17	CNS0155H	CNS0155H
23	68.8	4.5	963	17	CNS00A4L	CNS00A4L
24	68.4	4.5	1029	17	CNS01ZGM	CNS01ZGM
25	68	4.5	787	17	AQ740471	AQ740471
26	67.8	4.5	887	17	BF274559	BF274559
27	67.8	4.5	1101	17	CNS00U72	CNS00U72
28	67.6	4.4	876	17	CNS005G1	CNS005G1
29	67.6	4.4	1101	17	CNS00E07	CNS00E07
30	67.4	4.4	483	9	AU087226	AU087226
31	67.4	4.4	1101	17	CNS004ZM	CNS004ZM
32	67.4	4.4	1225	17	CNS0161D	CNS0161D
33	67	4.4	942	17	CNS018G5	CNS018G5
34	67	4.4	970	17	CNS0182A	CNS0182A
35	67	4.4	1092	17	CNS020K7	CNS020K7
36	67	4.4	1190	17	CNS020N7	CNS020N7
37	66.8	4.4	1027	17	CNS02750	CNS02750
38	66.6	4.4	928	17	CNS07ABZ	CNS07ABZ
39	66.6	4.4	928	17	CNS00DXY	CNS00DXY
40	66.6	4.4	1101	17	CNS000B8	CNS000B8
41	66.6	4.4	1101	17	CNS00EVL	CNS00EVL
42	66.6	4.4	1168	17	CNS06HDF	CNS06HDF
43	66.6	4.4	996	17	CNS00F0H	CNS00F0H
44	66.4	4.4	1135	17	CNS033GQ	CNS033GQ
45	66.4	4.4	1135	17	CNS033GQ	CNS033GQ

ALIGNMENTS

RESULT 1
LOCUS A1694972
DEFINITION we44h07.x1 NCI_CGAP_Lu24 Homo sapiens cDNA clone IMAGE:2343997 3', mRNA sequence.
ACCESSION A1694972
VERSION A1694972.1 GI:4982872
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 391)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapsb@mail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
CDNA Library Preparation: M. Bento Soares, Ph.D.
DNA Sequencing by: Greg Lennon, Ph.D.
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/ILMUT at: www.bio.lnl.gov/bbrp/image/image.html
Insert length: 672 Std Error: 0.00
Seq primer: -40UP from Gibco

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High quality sequence stop: 361.
Location/Qualifiers
1. 391
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:2343997"
/clone_lib="NCI CGAP Lu24"
/tissue_type="carcinoid"
/lab_host="DH108"
/note="Organ: lung; Vector: pTVT3D-Pac (Pharmacia) with a modified polylinker; Plasmid DNA from the normalized library NCI CGAP Lu5 was prepared and as circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneIDs 1414920-1417991 and 1520904-1522439). Subtraction by Bento Soares and M. Fatima Bonaldo."
BASE COUNT 66 a 123 c 96 g 106 t
ORIGIN

Query Match 24.9%; Score 378.2; DB 9; Length 391;
Best Local Similarity 99.2%; Pred. No. 3.3e-75;
Matches 380; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1138 TATTTGACACCTAGAAATTTTAGGCAAGTATAGTGGTATCATCTGCGGACCGCTTT 1197
Db 1 TTTTGGACCTAGAAATTTTAGGCAAGTATAGTGGTATCATCTGCGGACCGCTTT 60
QY 1198 TTTTCTTGACAGTAGTCTGTTGGGAGGAGTCTGCCACTCAGCTCTCTGCAGTCT 1257
Db 61 TTTTCTTGACAGTAGTCTGTTGGGAGGAGTCTGCCACTCAGCTCTCTGCAGTCT 120
QY 1258 CCGGCTCTCTCTGCGAGGATCGGTCAACGACCGCTCGCCGCTCTGCACCCAGCCAG 1317
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QY 1318 GTCCGACCTGCTCAGTCCGGTCTCAAGCTCAGCACCCTTTTATCCCGAGCAGC 1377
Db 181 GTCCGACCTGCTCAGTCCGGTCTCAAGCTCAGCACCCTTTTATCCCGAGCAGC 240
QY 1378 CTGATCGTCTGCTTCCCTCAGTCCGGAGCCACTGTAGTCCGACCGCGCTCTGA 1437
Db 241 CTGATCGTCTGCTTCCCTCAGTCCGGAGCCACTGTAGTCCGACCGCGCTCTGA 300
QY 1438 TATTCGGTGAAGTCTTCTCTGAGGTTTGGTCTCCGAGTCTCTGTGTAGTCCACCTT 1497
Db 301 TATTCGGTGAAGTCTTCTCTGAGGTTTGGTCTCCGAGTCTCTGTGTAGTCCACCTT 360
QY 1498 AGCGGTGTACGCTCTTTTGA 1520
Db 361 AGCGGTGTACGCTCTTTTGA 383

RESULT 2
AA324132 367 bp mRNA linear EST_20-APR-1997
LOCUS EST27025 Cerebellum II Homo sapiens cDNA 5' end, mRNA sequence.
DEFINITION AA324132.1 GI:1976479
ACCESSION
VERSION
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 367)
ADAMS, W.D., Kerlavage, A.R., Fleischmann, R.D., Fuldner, R.A., Bult, C.J., Lee, N.H., Kinkness, E.F., Weinstein, K.G., Gocayne, J.D., White, O., Sutton, G., Blake, J.A., Brandon, R.C., Mine, W.L., Cline, T.R., Cotton, M.D., Earle-Hughes, J., Fane, L.D., Fitzgerald, L.M., Fitzhugh, W.M., Fritchman, J.L., Geoghegan, N.S., Glodok, A., Gnehm, C.L., Hanna, M.C., Hedblom, E., Hinkle, P.S., Jr., Kelley, J.M., Kelley, J.C., Liu, L.-I., Marmaros, S.M., Merrick, J.M.,

Moreno-Palancas, R.F., McDonald, L.A., Nguyen, D.T., Pelligrino, S.M., Phillips, C.A., Ryder, S.E., Scott, J.L., Saudek, D.M., Shirley, R., Small, K.V., Spriggs, T.A., Utterback, T.R., Weidman, J.P., Li, Y., Bednarek, D.P., Cao, L., Cepeda, M.A., Coleman, T.A., Collins, E.J., Dimke, D., Feng, D.-P., Fertie, A., Fischer, C., Hastings, G.A., He, W.W., Hu, J.S., Greene, J.M., Gruber, J., Hudson, P., Kim, A.K., Kozak, D.L., Kunsch, C., Hung, J., Li, H., Meisner, P.S., Olsen, H., Raymond, L., Wei, Y.F., Wing, J., Xu, C., Yu, G.L., Ruben, S.M., Dillon, P.J., Pannone, M.R., Rosen, C.A., Haseltine, W.A., Fields, C., Fraser, C.M. and Venter, J.C.
Initial assessment of human gene diversity and expression patterns based upon 83 million nucleotides of cDNA sequence
NATURE 377 (6547 Suppl.), 3-174 (1995)
MEDLINE 96026280
COMMENT Contact: Kerlavage, AR
Bioinformatics
The Institute for Genomic Research
9712 Medical Center Drive, Rockville, MD 20850 USA
Tel: 3018699056
Fax: 3018699423
Email: arkerlav@tigr.org
For clone availability, additional sequence and expression information related to this EST, please check the TIGR Human Gene Index (<http://www.tigr.org/tldb/hgi/hgi.html>)
Seq primer: M13 Reverse.
Location/Qualifiers
1. 367
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/db_xref="taxon:9606"
/clone_lib="Cerebellum II"
/tissue_type="cerebellum"
/dev stage="adult"
/note="Organ: brain; Vector: pBluescript SK-; Site_1: EcoRI; Site_2: XhoI"
BASE COUNT 54 a 113 c 98 t 10 others
ORIGIN

Query Match 22.7%; Score 345.4; DB 9; Length 367;
Best Local Similarity 96.9%; Pred. No. 8.9e-68;
Matches 346; Conservative 0; Mismatches 11; Indels 0; Gaps 0;
QY 1164 GTGATAGTGGTATCATCTGCGGACCGCTTTTCTTTCGACGAGTCTGCTTCCG 1223
Db 1 GTGATAGTGGTATCATCTGCGGACCGCTTTTCTTTCGACGAGTCTGCTTCCG 60
QY 1224 GAGGAGTCTGCCACTCGAGCTCTCTGAGTCTCCGCTCTCTCTCAGGATCGGTCA 1283
Db 61 GAGGAGTCTGCCACTCGAGCTCTCTGAGTCTCCGCTCTCTCTCAGGATCGGTCA 120
QY 1284 ACGAGCGTCCGCGCTCTGACCCAGCGCTGCGCTCTCTCAGTCCGCTTCTC 1343
Db 121 ACGAGCGTCCGCGCTCTGACCCAGCGCTGCGCTCTCTCAGTCCGCTTCTC 180
QY 1344 AAGCCTCAGCACCCTCTTTTATCCCGAGCAGCTGAGTCTCTCAGTCCCGA 1403
Db 181 AAGCCTCAGCACCCTCTTTTATCCCGAGCAGCTGAGTCTCTCAGTCCCGA 240
QY 1404 CGCCACTGCTAGTCTGACCAACCGCGCTCTCTGATATTCGGTGA 1463
Db 241 CGCCACTGCTAGTCTGACCAACCGCGCTCTCTGATATTCGGTGA 300
QY 1464 GGTTGGTCTCCGATCTCTGTGTAGCCACCTTAGCGGTGTACGGTCTTTGAAA 1520
Db 301 GGTTGGTCTCCGATCTCTGTGTAGCCACCTTAGCGGTGTACGGTCTTTGAAA 357

RESULT 3
AL120587 474 bp mRNA linear EST 25-FEB-2000
LOCUS AL120587
DEFINITION DK229761E219 r1 761 (synonym: hamy2) Homo sapiens cDNA clone
VERSION DK229761E219 5', mRNA sequence.
ACCESSION AL120587

VERSION AL120587.1 GI:5926486
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 474)
AUTHORS Blum, H., Bauerbach, S., Mewes, H.W., Gassenhuber, J., and Wiemann, S.
TITLE EST (Blum, et al.)
JOURNAL Unpublished (1999)
COMMENT Contact: Blum H
MIPS

Am Klopferpitz 18a D-82152 Martinsried, Germany
This is the 5' sequence of the clone insert
Clone from S. Wiemann, Molecular Genome Analysis, German Cancer
Research Center (DKFZ); Email s.wiemann@dkfz-heidelberg.de;
sequenced by LMU (Ludwig Maximilians University,
Munich/Germany) within the cDNA sequencing consortium of the German
Genome Project.
a1 sequence also available.
This clone (DKFZ761E219) is available at the RZPD in Berlin.
Please contact the RZPD: Ressourcentzentrum, Heubnerweg 6, 14059
Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.
FEATURES
Location/Qualifiers
Source
1..474
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/clone="DKFZ761E219"
/clone_1="761 (synonym: hamy2)"
/tissue_type="amygdala"
/dev_stage="adult"
/lab_host="DH10B"
/note="Vector: pSport1; Site_1: NotI; Site_2: SalI"

BASE COUNT 84 a 138 c 141 g 111 t
ORIGIN
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Best Local Similarity 21.5%; Score 327; DB 9; Length 474;
Matches 327; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1194 GTTTTCTTCTGAGCAGTACTGCTTGGCGAGAGTGGCCCACTGAGCTCTTCTCA 1253
13 GTTTTCTTCTGAGCAGTACTGCTTGGCGAGAGTGGCCCACTGAGCTCTTCTCA 72
1254 GTTCCGGGCTCTCTCTGAGCAGTACTGCTTGGCGAGAGTGGCCCACTGAGCTCTTCTCA 1313
73 GTTCCGGGCTCTCTCTGAGCAGTACTGCTTGGCGAGAGTGGCCCACTGAGCTCTTCTCA 132
1314 CAGGTCGACACTGCTTCTGAGCAGTACTGCTTGGCGAGAGTGGCCCACTGAGCTCTTCTCA 1373
133 CAGGTCGACACTGCTTCTGAGCAGTACTGCTTGGCGAGAGTGGCCCACTGAGCTCTTCTCA 192
1374 CAGGTCGACACTGCTTCTGAGCAGTACTGCTTGGCGAGAGTGGCCCACTGAGCTCTTCTCA 1433
193 CAGGTCGACACTGCTTCTGAGCAGTACTGCTTGGCGAGAGTGGCCCACTGAGCTCTTCTCA 252
1434 CTGATATTTCTGAGCAGTACTGCTTGGCGAGAGTGGCCCACTGAGCTCTTCTCA 1493
253 CTGATATTTCTGAGCAGTACTGCTTGGCGAGAGTGGCCCACTGAGCTCTTCTCA 312
QY 1494 CTTAGGGGCTGAGCAGTCTTTGAAA 1520
Db 313 CTTAGGGGCTGAGCAGTCTTTGAAA 339

RESULT 4
BI457956 741 bp mRNA linear EST 21-AUG-2001
LOCUS 603198949F1 NIH_MGC_96 Homo sapiens cDNA clone IMAGE:5278356 5',
DEFINITION mRNA sequence.
ACCESSION BI457956
VERSION BI457956.1 GI:15248612
KEYWORDS EST.

SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 741)
AUTHORS NIH-MGC http://mgi.nci.nih.gov/
TITLE NIH-MGC
JOURNAL National Institutes of Health, Mammalian Gene Collection (MGC)
COMMENT Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cga@b-remail.nih.gov

Tissue Procurement: Miklos Palokovics, M.D., Ph.D.
cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
Toshitaki and Piero Carninci (RIKEN)
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: InCyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at:
http://image.llnl.gov
plate: L14M11703 row: a column: 13
High quality sequence stop: 741.
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Location/Qualifiers
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/clone_1="NIH_MGC_96"
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/note="Organ: brain; Vector: pBluescript (modified
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size-selected for average insert size 2.3 kb and
normalized to ROT 5. This is a primary library enriched
for full-length clones and constructed using the
cap-trapper method (Carninci, in preparation). Library
constructed by M. Brownstein (NIH/NHGRI, National
Institutes of Health). Note: this is a NIH_MGC Library."

BASE COUNT 177 a 180 c 218 g 166 t
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Best Local Similarity 19.7%; Score 299.2; DB 13; Length 741;
Matches 301; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

1217 GCTTGGGAGAGAGTCTCCCACTGACCTCTTGGCAGTCTCTCTGACAGA 1276
6 GCTTGGGAGAGAGTCTCCCACTGACCTCTTGGCAGTCTCTCTGACAGA 65
QY 1277 TCGTCAACGACGCGTGGCGGCTCTGACAGCCAGGTCGACCTTCACTCC 1336
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1337 GATTCTCAAGGCTCAGACCATTTTATCCCGAGACCTGATTCGTCCTCA 1396
126 GATTCTCAAGGCTCAGACCATTTTATCCCGAGACCTGATTCGTCCTCA 185
QY 1397 GTCCGAGCCCACTGATTCGACCAACGCGGCTTGTGATTTTCGATCTTTC 1456
186 GTCCGAGCCCACTGATTCGACCAACGCGGCTTGTGATTTTCGATCTTTC 245
QY 1457 CTGAGAGTTTGGTCTCCGATCTCTGATGAGCAGCTTGAAGGATCTTTG 1516
246 CTGAGAGTTTGGTCTCCGATCTCTGATGAGCAGCTTGAAGGATCTTTG 305
QY 1517 AAAA 1520
Db 306 AAAA 309

RESULT 5
BG714610 785 bp mRNA linear EST 08-MAY-2001
LOCUS BG714610
DEFINITION 602676733F1 NIH_MGC_96 Homo sapiens cDNA clone IMAGE:4799121 5',

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BG716320
LOCUS
DEFINITION
  BG716320.1 GI:13993541
  mRNA sequence.
  EST.
  human.
  Homo sapiens
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  Eukaryota; Eutheria; Primates; Catarrhini; Hominidae; Homo.
  NIH-MGC http://mgi.nci.nih.gov/.
  National Institutes of Health, Mammalian Gene Collection (MGC)
  Unpublished (1999)
  Contact: Robert Strausberg, Ph.D.
  Email: cgaabs-remail.nih.gov
  Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
  cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
  Toshiyuki and Piero Carninci (RIKEN)
  cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
  DNA Sequencing by: Incyte Genomics, Inc.
  Clone distribution: MGC clone distribution information can be
  found through the I.M.A.G.E. Consortium/LLNL at:
  http://image.llnl.gov
  Plate: L1AM10688 row: a column: 10
  High quality sequence stop: 777.
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    /clone_lib="NIH_MGC_96"
    /tissue_type="hypothalamus"
    /lab_host="DH10B"
    /note="Organ: brain; Vector: pBluescriptR (modified
    pBluescript KS+); Site 1: BamHI; Site 2: SalI-XhoI (gtcgag
    ); Oligo-dT primed using primer 5'-TTTTTTTTTTTNN-3',
    size-selected for average insert size 2.3 kb and
    normalized to R0T 5. This is a primary library enriched
    for full-length clones and constructed using the
    Cap-trapper method (Carninci, in preparation). Library
    constructed by M. Brownstein (NHGRI, National
    Institutes of Health). Note: this is a NIH_MGC Library."
  BASE COUNT      196 a 188 c 224 g 177 t
  ORIGIN
    Query Match      19.7%; Score 299.2; DB 12; Length 785;
    Best Local Similarity 99.0%; Pred. No. 2.6e-57; Indels 0; Gaps 0;
    Matches 301; Conservative
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  DB 66 TCGGTCAACGAGCGTCTGCCACCTGCGAGCTCTGCGAGTCTCGGCTCTCTCTCGAGTC 125
  QY 1337 GGTTCCTAAAGCCTCAGACCATCTTTATCCCGAGCAGCTCGATCGTCTCGATCTCA 1396
  DB 126 GGTTCCTAAAGCCTCAGACCATCTTTATCCCGAGCAGCTCGATCGTCTCGATCTCA 185
  QY 1397 GTCCGAGCGCCACTGCTAGGTCTGAGTCTGATATTTTGGTGGTCTTTTC 1456
  DB 186 GTCCGAGCGCCACTGCTAGGTCTGAGTCTGATATTTTGGTGGTCTTTTC 245
  QY 1457 CTGTGAGGTTGGTCTCCCGATCTCTGTGTAGCCACCTTAGGCGTGTAGCGTCTTTG 1516
  DB 246 CTGTGAGGTTGGTCTCCCGATCTCTGTGTAGCCACCTTAGGCGTGTAGCGTCTTTG 305
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  human.
  Homo sapiens
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  Eukaryota; Eutheria; Primates; Catarrhini; Hominidae; Homo.
  NIH-MGC http://mgi.nci.nih.gov/.
  National Institutes of Health, Mammalian Gene Collection (MGC)
  Unpublished (1999)
  Contact: Robert Strausberg, Ph.D.
  Email: cgaabs-remail.nih.gov
  Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
  cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
  Toshiyuki and Piero Carninci (RIKEN)
  cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
  DNA Sequencing by: Incyte Genomics, Inc.
  Clone distribution: MGC clone distribution information can be
  found through the I.M.A.G.E. Consortium/LLNL at:
  http://image.llnl.gov
  Plate: L1AM10687 row: j column: 21
  High quality sequence stop: 713.
  Location/Qualifiers
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    pBluescript KS+); Site 1: BamHI; Site 2: SalI-XhoI (gtcgag
    ); Oligo-dT primed using primer 5'-TTTTTTTTTTTNN-3',
    size-selected for average insert size 2.3 kb and
    normalized to R0T 5. This is a primary library enriched
    for full-length clones and constructed using the
    Cap-trapper method (Carninci, in preparation). Library
    constructed by M. Brownstein (NHGRI, National
    Institutes of Health). Note: this is a NIH_MGC Library."
  BASE COUNT      195 a 195 c 229 g 179 t
  ORIGIN
    Query Match      19.7%; Score 299.2; DB 12; Length 798;
    Best Local Similarity 99.0%; Pred. No. 2.6e-57; Indels 0; Gaps 0;
    Matches 301; Conservative
  QY 1217 GCTTGGGAGGAGTCTGCCACCTGCGAGCTCTGCGAGTCTCGGCTCTCTCTCGAGGA 1276
  DB 6 GCTTGGGAGGAGTCTGCCACCTGCGAGCTCTGCGAGTCTCGGCTCTCTCTCGAGGA 65
  QY 1277 TCGGTCAACGAGCGTCTGCCACCTGCGAGCTCTGCGAGTCTCGGCTCTCTCTCGAGTC 1336
  DB 66 TCGGTCAACGAGCGTCTGCCACCTGCGAGCTCTGCGAGTCTCGGCTCTCTCTCGAGTC 125
  QY 1337 GGTTCCTAAAGCCTCAGACCATCTTTATCCCGAGCAGCTCGATCGTCTCGATCTCA 1396
  DB 126 GGTTCCTAAAGCCTCAGACCATCTTTATCCCGAGCAGCTCGATCGTCTCGATCTCA 185
  QY 1397 GTCCGAGCGCCACTGCTAGGTCTGAGTCTGATATTTTGGTGGTCTTTTC 1456
  DB 186 GTCCGAGCGCCACTGCTAGGTCTGAGTCTGATATTTTGGTGGTCTTTTC 245
  QY 1457 CTGTGAGGTTGGTCTCCCGATCTCTGTGTAGCCACCTTAGGCGTGTAGCGTCTTTG 1516
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  QY 1517 AAAA 1520
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Db 306 AAAA 309

RESULT 7
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LOCUS BI602678
DEFINITION 603252017F1 NIH_MGC_96 Homo sapiens cDNA clone IMAGE:5303692 5',
mRNA sequence.
ACCESSION BI602678
VERSION BI602678.1 GI:15495617
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 586)
NIH-MGC http://mgs.nci.nih.gov/
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgabbs-remail.nih.gov
Tissue Procurement: Miklos Palokovits, M.D., Ph.D.
CDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiroki
Toshiyuki and Piero Carninci (RIKEN)
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (ILNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/ILNL at:
http://image.llnl.gov
Plate: LLAM11769 row: a column: 05
High quality sequence stop: 586.
Location/Qualifiers
1. 586
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/db_xref="taxon:9606"
/clone="IMAGE:5303692"
/clone_1ib="NIH_MGC_96"
/issue_type="hypothalamus"
/lab_host="DH10B"
/note="Organ: brain; Vector: pBluescript (modified
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); Oligo-dt primed using primer 5'-TTTTTTTTTTTTTTVN-3',
size-selected for average insert size 2.3 kb and
normalized to 10⁵. This is a primary library enriched
for full-length clones and constructed using the
Cap-trapper method (Carninci, in preparation). Library
constructed by M. Brownstein (NIH/NHGRI, National
Institutes of Health). Note: this is a NIH_MGC Library."

FEATURES
source

BASE COUNT 129 a 148 c 182 g 127 t

Query Match 17.2%; Score 261.6; DB 13; Length 586;
Best Local Similarity 98.5%; Pred. No. 8.6e-49;
Matches 264; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

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1 ACGGCGCGCTCTCTCCGAGATGCGTCAACGACCGTGGCGGCTCTGACCCAG 60
QY 1313 CCGAGGTGCGCACTGCTTCAAGTCTCAAGGCTCAGACACATCTTTATCCCGA 1372
61 CACAGGTGCGCACTGCTTCAAGTCTCAAGGCTCAGACACATCTTTATCCCGA 120
QY 1373 GAGGCTGAGTCTGCTTCCGTCAGTCCGAGCGGCACTGCTAGGTCGACGCGCGCT 1432
Db 121 GAGGCTGAGTCTGCTTCCGTCAGTCCGAGCGGCACTGCTAGGTCGACGCGCGCT 180
QY 1433 TCTGATATTTGCGTGAAGTCTTTCTGAGAGGTTGGTCTCCCGATCTCTGTGTAGCC 1492
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Db 241 ACCTTAGCGGTAGGCTCTTTGAAAA 268

RESULT 8
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LOCUS B0601726
DEFINITION MI-P-HO-afw-d-02-1-UM s1 MI-P-HO Sus scrofa cDNA clone
MI-P-HO-afw-d-02-1-UM 3', mRNA sequence.
ACCESSION B0601726
VERSION B0601726.1 GI:21548452
KEYWORDS EST.
SOURCE pig.
ORGANISM Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suidae; Sus.
1 (bases 1 to 459)
Bonaldo, M.F., Lennon, G. and Soares, M.B.
Normalization and subtraction: two approaches to facilitate gene
discovery
Genome Res. 6 (9), 791-806 (1996)
Contact: Tugue CK
97044477
Molecular Genetics Laboratory, Department of Animal Science
Iowa State University
201 Kildee Hall, Ames, IA 50011-3150, USA
Tel: 5152944252
Fax: 5152942401
Email: ctugue@iastate.edu
Tissue Procurement: Dr. Chris Tugue, Iowa State University
CDNA Library Preparation: Dr. M. Bento Soares, University of Iowa
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Researchers may obtain clones from Research
Genetics (www.resgen.com).
Seq primer: M13 FORWARD
POLYA=No.

FEATURES
source

Location/Qualifiers
1. 459
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/db_xref="taxon:9823"
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/clone_1ib="MI-P-HO"
/lab_host="DH10B (Life Technologies)"
/note="Vector: pT73D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: BcoRI; The MI-P-HO
library is a normalized library derived from hypothalamus
at estrus days 0 and 12 and ovary at estrus days 0, 5 and
12. For a detailed description of the library from which
this clone was derived, please visit our web site at
http://pigdb.genome.iastate.edu/
TAG_L1B=MI-P-HO
TAG_TISSUE=d_0 hypothalamus
TAG_SEQ=TAGATG"
BASE COUNT 79 a 152 c 126 g 102 t

Query Match 10.8%; Score 164.4; DB 14; Length 459;
Best Local Similarity 74.0%; Pred. No. 8.8e-27;
Matches 256; Conservative 0; Mismatches 71; Indels 19; Gaps 3;

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1 GTTTTTTCTTGAGAGTAGCTGCTGCTGAGGAGGAGTCTGCACTGCGCTCTGTC 60
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Db 121 GCCGAGCAACGCGCTGCTGTCGCTTTTGTATCTTCAAGACACCTTCTCTCTGCA 180

[illegible]

GenCore version 5.1.3
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 8, 2003, 16:37:35 / Search time 351 Seconds
(without alignments)
9752.243 Million cell updates/sec

Title: US-09-847-665-4
Perfect score: 1520

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Scoring table:

IDENTITY_NUC
Gapop 10.0, Gapext 1.0

Number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1520	100.0	1520	24	ABK10803
2	1520	100.0	3699	24	ABK10805
3	522.4	34.4	2720	24	ABK10802
4	380.6	25.0	629	21	AA01166
5	233.8	15.4	1725	24	ABK10801
6	166.8	11.0	2819	24	ABK10806
7	67	4.4	513445	22	AA161373
8	65.2	4.3	17967	24	AB131015
9	64.8	4.3	6419	24	ABL32267

10	64.4	4.2	19124	18	AA72882	Plasmodium var-7 g
11	64.4	4.2	19124	21	AA298287	Plasmodium var-7 p
12	63.8	4.2	83391	24	AB067093	Human angiotensin
13	63	4.1	3991	22	AA016633	Human novel protei
14	63	4.1	6286	22	AA546591	Tumour suppressor
15	63	4.1	17234	24	AB067018	Human angiotensin
16	63	4.1	18817	24	ABL70162	Chemically treated
17	63	4.1	18817	24	ABL70162	Human metastasis a
18	62.8	4.1	7849	24	AA022278	Chemically treated
19	62.8	4.1	7849	24	AA022278	Human immune syste
20	62.6	4.1	7597	24	AB133013	Human immune syste
21	62.6	4.1	16217	24	AB132634	Human immune syste
22	62.4	4.1	6109	24	AB132226	Human immune syste
23	62.4	4.1	6109	24	AB132226	Human immune syste
24	62.4	4.1	7814	24	AA546530	Human gene regulat
25	62.2	4.1	6317	24	AA546530	Tumour suppressor
26	62.2	4.1	6317	24	AB149311	Human polynucleoti
27	62.2	4.1	5875	24	AB132408	Human immune syste
28	61.8	4.1	17594	24	AB132408	Human immune syste
29	61.4	4.0	11092	24	AB134026	Human immune syste
30	61	4.0	7346	24	AB133513	Human immune syste
31	60.8	4.0	6175	24	AB133507	Human immune syste
32	60.8	4.0	37515	24	AB133507	Human immune syste
33	60.6	4.0	6082	24	AB066998	Human angiotensin
34	60.6	4.0	7862	24	AB133753	Human immune syste
35	60.6	4.0	8085	22	AA546480	Human immune syste
36	60.6	4.0	8085	22	AA546480	Tumour suppressor
37	60.6	4.0	16173	24	AB133987	Human DNA for stag
38	60.4	4.0	5267	24	AB067043	Human metastasis a
39	60.4	4.0	6048	24	AB067002	Human angiotensin
40	60.4	4.0	6306	22	AA545516	Human angiotensin
41	60.4	4.0	6306	24	ABK28458	Chemically treated
42	60.4	4.0	6370	24	ABL70567	DNA transcription
43	60.4	4.0	6370	24	ABK31348	Chemically treated
44	60.2	4.0	4590	7	AA060472	Signal transductio
45	60.2	4.0	6027	24	ABN80172	Sequence encoding

ALIGNMENTS

RESULT 1	ABK10803	standard; DNA; 1520 BP.
ID	ABK10803	
XX	ABK10803;	
AC		
XX		
XX	21-MAY-2002 (first entry)	
DT		
XX		
XX	Human nucleosomal assembly protein IL2 (NAPIL2 or BPX) promoter.	
DE		
XX		
XX	Human nucleosomal assembly protein IL2; NAPIL2; BPX; nootropic;	
KW	neurotropic; cytoskeletal; neural system defect; cancer;	
KW	neuronal cell tumour; gene therapy; neurodegenerative disease;	
KW	Parkinson's disease; Alzheimer's disease; spina bifida; anencephaly;	
KW	brain structure; brain organisation; X-chromosome linked disorder;	
KW	nucleosome activity; cell cycle; cyclin; transcription factor control;	
XX	histone; promoter; ds.	
XX		
OS	Homo sapiens.	
FN		
XX	WO200185995-A2.	
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PD	15-NOV-2001.	
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PF	04-MAY-2001; 2001WO-1B00960.	
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XX	05-MAY-2000; 2000US-202111P.	
PR		
XX	(INSP) INST PASTEUR.	
PA	Soybean 318013 reg	
XX	(CNRS) CNRS CENT NAT RECH SCI.	
XX		
PI	Avner P. Rogner UC, Spyropoulos D, Rougeulle C;	

Thu Jan 9 09:34:26 2003

XX WPI; 2002-075254/10.
XX Screening neural system defects in mammal, comprises detecting a
XX modification of Napi12 and/or Napi12 gene chromosomal material, that
XX causes a loss of biological function and correlating the modification
XX with the defect
XX
XX Claim 24; Fig 10; 83pp; English.
XX The invention describes a method of screening mammalian neural system
XX defects, comprising detecting a modification of the nucleosomal assembly
XX protein Napi12 and/or Napi12 gene in a human and/or mouse, where
XX modification is substitution, deletion, frame-shift, insertion aberrant
XX or altered epigenetic control, that causes a loss of function in the gene
XX and correlating the gene modification with a potential for a neural
XX system defect. The invention includes a neuronal cell containing a
XX recombinant chromosome on which Napi12 is located. The cell is useful for
XX screening therapeutic compounds by introducing a compound to be screened
XX and correlating change in the proliferation of cells with the activity of
XX the compound, where change in proliferating cells is preferably a control
XX of cancer of neural cells. The Napi12 promoter is useful for predicting
XX of cancer of neural origin, and for genetic therapy of neurodegenerative
XX diseases such as Parkinson's, Alzheimer's diseases or accidents. Napi12
XX is also implicated in spina bifida with or without anencephaly, loss of
XX brain structure/organisation, X-chromosome linked disorders and
XX inappropriate control of nucleosome activity in neurons, neuronal cell
XX cycle, cyclins, DNA binding transcription factors, histones and histone
XX shuttling. This sequence represents the promoter of the human
XX nucleosomal assembly protein (Napi12 or BPX) gene, a homologue of the
XX murine nucleosomal assembly protein (Napi12 or Bpx) gene described in
XX the invention.
XX
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XX Query Match 100.0%; Score 1520; DB 24; Length 1520;
XX Best Local Similarity 100.0%; Pred. No. 0;
XX Matches 1520; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 841 CGACATTTTATAGTTTACGTTTCTCTGAGCTCTCTGGAAGCAATAAAGTATATATCTGTT 900
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RESULT 2
ABK10805

ID	ABK10805 standard; DNA; 3699 BP.
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AC	ABK10805;
XX	
DT	21-MAY-2002 (first entry)
DE	DNA encoding human nucleosomal assembly protein (NAPIL2/BPX) protein.
KW	Human nucleosomal assembly protein; NAPIL2; BPX; nucleotropic;
KW	neurotropic; cytotostatic; neural system defect; cancer;
KW	neuronal cell tumour; gene therapy; neurodegenerative disease;
KW	Parkinson's disease; Alzheimer's disease; spina bifida; anencephaly;
KW	brain structure; brain organisation; X-chromosome linked disorder;
KW	nucleosome activity; cell cycle; cyclin; transcription factor control;
KW	histone; gene; ds.
OS	Homo sapiens.
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PF	04-MAY-2001; 2001MO-IB09960.
PR	05-MAY-2000; 2000US-202111P.
PA	(INSP) INST PASTEUR.
PI	(CNRS) CNRS CENT NAT RECH SCI.
PT	Ayner P, Rogner UC, Spyropoulos D, Rougeulle C,
DR	WPI; 2002-075254/10.
XX	
PT	Screening neural system defects in mammal, comprises detecting a
PT	modification of NAPIL2 and/or Napil2 gene chromosomal material, that
PT	causes a loss of biological function and correlating the modification
XX	with the defect -
XX	
PS	Disclosure; Fig 13; 83pp; English.
XX	
CC	The invention describes a method of screening mammalian neural system
CC	defects, comprising detecting a modification of the nucleosomal assembly
CC	protein NAPIL2 and/or Napil2 gene in a human and/or mouse, where
CC	modification is substitution, deletion, frame-shift, insertion aberrant
CC	or altered epigenetic control, that causes a loss of function in the gene
CC	and correlating the gene modification with a potential for a neural
CC	system defect. The invention includes a neuronal cell containing a
CC	recombinant chromosome on which Napil2 is located. The cell is useful for
CC	screening therapeutic compounds by introducing a compound to be screened
CC	and correlating change in the proliferation of cells with the activity of
CC	the compound, where change in proliferating cells is preferably a control
CC	of cancer of neural origin. The Napil2 promoter is useful for predicting
CC	tumour of neural cells. The Napil2 promoter is useful for predicting
CC	diseases such as Parkinson's, Alzheimer's diseases or accidents. Napil2
CC	is also implicated in spina bifida with or without anencephaly, loss of
CC	brain structure/organisation, X-chromosome linked disorders and
CC	inappropriate control of nucleosome activity in neurons, neuronal cell
CC	cycle, cyclins, DNA binding transcription factors, histones and histone
CC	shuttling. This sequence represents the human nucleosomal assembly
CC	protein (NAPIL2 or BPX) gene, a homologue of the murine nucleosomal
CC	assembly protein (Napil2 or Bpx) gene described in the invention.
XX	
SQ	Sequence 3699 BP; 1144 A; 636 C; 805 G; 1114 T; 0 other;
Query Match	100.0%; Score 1520; DB 24; Length 3699;
Best Local Similarity	100.0%; Pred. No. 0;
Matches 1520; Conservative	0; Mismatches 0; Indels 0; Gaps 0;

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Qy	61	AAACAGTTTTAATCTGTGATAGTAACAATCTTAAATCTGGAAAAATATATGTCCT	12
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Qy	181	AACTTACCAATATATTAAGTATAGAGAGAAATTTTAATATCTGCAAAAGCTTCATCTTA	24
Db	181	AACTTACCAATATATTAAGTATAGAGAGAAATTTTAATATCTGCAAAAGCTTCATCTTA	24
Qy	241	TAAATACATATATCAAAATGTTTAAACAATTCCTTAAATGCTGAGATTAGATTATTCGAA	30
Db	241	TAAATACATATATCAAAATGTTTAAACAATTCCTTAAATGCTGAGATTAGATTATTCGAA	30
Qy	301	TTAACTCAAAAGCATCAGCAAAATGTTATGATTTCTAATAATAAACATATACCTTCATTT	36
Db	301	TTAACTCAAAAGCATCAGCAAAATGTTATGATTTCTAATAATAAACATATACCTTCATTT	36
Qy	361	TGGCTTTTGTATATATATATATTAATTTCTAAGGCGCTTAAAGCCAGATTAGAGAGAG	42
Db	361	TGGCTTTTGTATATATATATATTAATTTCTAAGGCGCTTAAAGCCAGATTAGAGAGAG	42
Qy	421	CAGAAAGCATATGAGACTGGGGTATTTTAAGCCAGCAACTGGTTATATGTGGTT	48
Db	421	CAGAAAGCATATGAGACTGGGGTATTTTAAGCCAGCAACTGGTTATATGTGGTT	48
Qy	481	AATGTCTGATATGTTTACTAGTCAGATGATGTGTAACAACAATCTAGTTTTTCATACA	54
Db	481	AATGTCTGATATGTTTACTAGTCAGATGATGTGTAACAACAATCTAGTTTTTCATACA	54
Qy	541	GGCCCTCATTCGCCCCCACTGCGATCGGACTTCTCTCCCTCCCTCACAAGAAATGTTT	60
Db	541	GGCCCTCATTCGCCCCCACTGCGATCGGACTTCTCTCCCTCCCTCACAAGAAATGTTT	60
Qy	601	CGAGAAATTTTCAACCTTAATCATATAGCTTGTGAAAAATATCCGCAAAACATATATAG	66
Db	601	CGAGAAATTTTCAACCTTAATCATATAGCTTGTGAAAAATATCCGCAAAACATATATAG	66
Qy	661	AATATTTAAATATACGACAGCCCACTTAAAGACCATCATGCTAATTCCTGTGTTTTTA	72
Db	661	AATATTTAAATATACGACAGCCCACTTAAAGACCATCATGCTAATTCCTGTGTTTTTA	72
Qy	721	ATCTTTGAAGGCTTTGTTTATCAGCTCTTCAACATCCACTCTCCCTCCCAAGCTCC	78
Db	721	ATCTTTGAAGGCTTTGTTTATCAGCTCTTCAACATCCACTCTCCCTCCCAAGCTCC	78
Qy	781	CGATCTAAATATCAAAAGAGATGATTAAGATGGGTGGTGCCTTGTCTTCTCATTTGT	84
Db	781	CGATCTAAATATCAAAAGAGATGATTAAGATGGGTGGTGCCTTGTCTTCTCATTTGT	84
Qy	841	CGACATTTTACTTTCGTTTTCTGTGACTCTCTGAAAGCATTAAGATTAATATCTGTT	90
Db	841	CGACATTTTACTTTCGTTTTCTGTGACTCTCTGAAAGCATTAAGATTAATATCTGTT	90
Qy	901	AAAAGTTGATGATGACATTAATGAACGCAATGGGATTCGAAAAATCTCTGGGAGATG	96
Db	901	AAAAGTTGATGATGACATTAATGAACGCAATGGGATTCGAAAAATCTCTGGGAGATG	96
Qy	961	GGCTAGAGAGACGAGAGAGAGTGTGATGAATCAGCCATTTAGAAAGCTGGGAAAGGTAG	102
Db	961	GGCTAGAGAGACGAGAGAGAGTGTGATGAATCAGCCATTTAGAAAGCTGGGAAAGGTAG	102
Qy	1021	CAGAGTTGAAAACCTTGATGATCTATATATTTACCTGGGCTCGGGTTTGTACATCTACA	1080
Db	1021	CAGAGTTGAAAACCTTGATGATCTATATATTTACTGGCTCTGGGTTTGTACAGCTCTACA	1080

RESULT 4

ID AAC01166 standard; cDNA; 629 BP.

AC AAC01166;

DT 06-OCT-2000 (first entry)

DE Human secreted protein 5' EST, SEQ ID NO: 1164.

KM Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation; gene therapy; chromosome mapping; 85.

OS Homo sapiens.

PN EPI033401-A2.

PD 06-SEP-2000

PR 21-FEB-2000; 2000EP-0200610.

PR 26-FEB-1999; 99US-0122487.

PA (GEST) GENSET.

PI Dumas Milne Edwards J, Duclert A, Giordano J;

DR WPI; 2000-500381/45.

DR P-PSDB; AAG01160.

PT New nucleic acid that is a 5' expressed sequence tag (5' EST) for

PT diagnosing, forensic, gene therapy and chromosome mapping procedures -

PS Claim 1; SEQ ID 1164; 71bp + CD-ROM; English.

XX The present sequence is one of a large number of 5' ESTs derived from
CC mRNAs encoding secreted proteins. An ORF has been identified within the
CC sequence. The 5' ESTs were prepared from total human RNA or polyA+ RNAs
CC derived from 30 different tissues. EST sequences usually correspond
CC mainly to the 3' untranslated region (UTR) of the mRNA because they are
CC often obtained from oligo-dT primed cDNA libraries. Such ESTs are not
CC mRNAs and even in those cases where longer cDNA sequences have been
CC obtained, the full 5' UTR is rarely included. 5' ESTs are derived from
CC mRNAs with intact 5' ends and can therefore be used to obtain full length
CC cDNAs and genomic DNAs. 5' ESTs are also used in diagnostic, forensic,
CC gene therapy and chromosome mapping procedures. They are used to obtain
CC upstream regulatory sequences and to design expression and secretion
CC vectors.

XX SQ Sequence 629 BP; 128 A; 162 C; 191 G; 143 T; 5 other;

XX Query Match 25.0%; Score 380.6; DB 21; Length 629;

XX Best Local Similarity 99.7%; Pred. No. 6.1e-78;

XX Matches 380; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1140 TTTCGACCTAGAAATTTTGGCAAGATGATGCTGCGTAATCACTGCGACCGTTT 1199

DB 3 TTTCGACCTAGAAATTTTGGCAAGATGATGCTGCGTAATCACTGCGACCGTTT 62

QY 1200 TTTCGACCTAGAAATTTTGGCAAGATGATGCTGCGTAATCACTGCGACCGTTT 1259

DB 63 TTTCGACCTAGAAATTTTGGCAAGATGATGCTGCGTAATCACTGCGACCGTTT 122

QY 1260 GGCTCTCTGAGAGATCGGTCAACGACGCGCTCGGCTCTGACCCAGCCAGGT 1319

DB 123 GGCTCTCTGAGAGATCGGTCAACGACGCGCTCGGCTCTGACCCAGCCAGGT 182

QY 1320 CGCAGCTCTGAGAGATCGGTCAACGACGCGCTCGGCTCTGACCCAGCCAGGT 1379

DB 183 CGCAGCTCTGAGAGATCGGTCAACGACGCGCTCGGCTCTGACCCAGCCAGGT 242

QY 1380 GATTCGTCCTTCCTCAGTCCGAGACGCACTGATGTCGACCAACCGCCGCTTGTATA 1439

DB 243 GATTCGTCCTTCCTCAGTCCGAGACGCACTGATGTCGACCAACCGCCGCTTGTATA 302

QY 1440 TTTCGAGTGTCTTTTCCCTGTCGAGGTTTGGTCTCCGATCTGTGTAGCCACCTTNG 1499

DB 303 TTTCGAGTGTCTTTTCCCTGTCGAGGTTTGGTCTCCGATCTGTGTAGCCACCTTNG 362

QY 1500 GCGTGTACGTCCTTTGAAAA 1520

DB 363 GCGTGTACGTCCTTTGAAAA 383

RESULT 5

ID ABK10360 standard; DNA; 1725 BP.

AC ABK10360;

DT 21-MAY-2002 (first entry)

DE Murine nucleosomal assembly protein IL2 (Nap1L2 or Bpx) promoter.

KM Murine nucleosomal assembly protein IL2; Nap1L2; Bpx; noctropic;

KM neurotropic; cytoskeletal; neural system defect; cancer;

KM Parkinson's disease; Alzheimer's disease; spina bifida; anencephaly;

KM brain structure; brain organization; X-chromosome linked disorder;

KM histone; promoter; ds; mouse.

OS Mus sp.

PN WO200185995-A2.

PD 15-NOV-2001.

PF 04-MAY-2001; 2001WO-1B00960.

PR 05-MAY-2000; 2000US-202111P.

PA (INSP) INST PASTEUR.

PA (CNRS) CNRS CENT NAT RECH SCI.

PI Aymer P, Rogner UC, Spyropoulos D, Rougeulle C;

DR WPI; 2002-075254/10.

PT Screening neural system defects in mammal, comprises detecting a

PT modification of Nap1L2 and/or Nap1L2 gene chromosomal material, that

PT causes a loss of biological function and correlating the modification

PT with the defect -

PS Claim 32; Fig 7; 83pp; English.

XX The invention describes a method of screening mammalian neural system
CC defects, comprising detecting a modification of the nucleosomal assembly
CC protein Nap1L2 and/or Nap1L2 gene in a human and/or mouse, where
CC modification is substitution, deletion, frame-shift, insertion aberrant
CC or altered epigenetic control, that causes a loss of function in the gene
CC system defect. The invention includes a neuronal cell containing a
CC recombinant chromosome on which Nap1L2 is located. The cell is useful for
CC screening therapeutic compounds by introducing a compound to be screened
CC and correlating change in the proliferation of cells with the activity of
CC the compound, where change in proliferation of cells is preferably a control
CC of cancer of neural cells. The Nap1L2 promoter is useful for predicting
CC tumour of neural origin, and for genetic therapy of neurodegenerative
CC diseases such as Parkinson's, Alzheimer's diseases or accidents. Nap1L2
CC is also implicated in spina bifida with or without anencephaly, loss of
CC brain structure/organisation, X-chromosome linked disorders and
CC inappropriate control of nucleosome activity in neurons, neuronal cell
CC cycle, cyclins, DNA binding transcription factors, histones and histone

Thu Jan 9 09:34:26 2003

CC shuttling. This sequence represents a murine nucleosomal assembly protein
CC IL2 (NapIL2 or Bpx) promoter fragment, described in the invention.
XX
SQ

Sequence 1725 BP; 428 A; 443 C; 391 G; 463 T; 0 other;
Query Match 15.4%; Score 233.8; DB 24; Length 1725;
Best Local Similarity 65.1%; Pred. No. 5.4e-44;
Matches 486; Conservative 0; Mismatches 222; Indels 39; Gaps 8;
QY 580 CTCCCTCACAGGAATGTTTCGAGAAATTTTCAACCTAAATCATATATAGCTTGTGAAAA 639
Db 74 CTTCTCATCAGGAATATTATGAGAAATTTTCCCATTTTAAATCACACAGGTTGTGAAAA 133
QY 640 ATACGCAAAACATATATAGAAATTTTAAATTAATCACTGAC----ACGCCACCTAAAGACCA 695
Db 134 TTACAGA-AACCAAGGTACAGAAATTTTAAACCACTGTCAAGTTACATCATCAAGGCCA 192
QY 696 TCAGTGCTAATTCCTGGTGTGTTTAAATCTTTGAAAGCTTTTATCAGCTCTTCCACCA 755
Db 193 CTAATGCTTATTTTGGTAAATTTTAAACCTCAAGGATCTCTTTGGGCTCTCTCACTA 252
QY 756 TCCACCTCTCCCTCCAGGTCCCGATCTAAATCAAGAGATTTGATTAGATGGGT 815
Db 253 CCCTCTCTCT--CTTCCAGAGCCTCAGGTTATACCAAGGATAGACTAAAGACAATC 310
QY 816 GGGTGGCTTGTCTCTCTCATTTGTCGACATTTAGTTAGCTTTTCTCTGAGCTCTCTGG 875
Db 311 CAGTACTTGGCCATTTTTCATTC-----CTTGTCACTGTTCCATATAGCTCTTTTG 365
QY 876 AAAGCAATAAGTATATATCTGTTTAAAGTTGGATGAATGAATCAATGAACGCAATGGG 935
Db 366 AAATATGAACATATATAGTATCAGTTGAAACCGAATGAATGA-----TACTGCA 414
QY 936 ATCCGAAACCTCTCGGAGATGGGCTAGAGGAGGAGGAGGTGGATGAATCAGCC 995
Db 415 TTTCTGCAAAATCCACAGCTATAGGTGGAAGATGAGCCATAGGTGGAGGATCAGCC 474
QY 996 ATGTTAGAGGCTGGGAAGGTGACGAGCTTGAACAACTTGTAGATCTTAATTAATTTACT 1055
Db 475 ATATTAGAAATCTGGGAAGCAAGAGGTGTGAAATTTGATTCTACTTAATTTACT 534
QY 1056 GGCTCTGGTTTGTGAGTCACTATATGCAAGCAAAATGAGATTAGACATGTTGTGGAG 1115
Db 535 GCTCAGGATTTGTCATCACTGAGCTGCAATGAGATGAGATGAGAGAGTCTCTGGAG 594
QY 1116 GGAAGGAGTGCAGCAATCTATTGCACT--AGAAATTTTGGCAAGTATGATCTGC 1174
Db 595 GGAAGG-GGTGACGAGCAACCTGCATACCTTAAATAAAGAGGCTGAGAGACAATGC 653
QY 1175 GTAAATCACTGCGGCACCGTTTTCCTT-----GACGAGTACTGCTT 1220
Db 654 GTAAATCACTGCGGCACCGTTTTCCTTCCATCCCTCCGCCCCGAGTGGTGGAGCAGCTG 713
QY 1221 GCGGAGGAGTGTGCGCACTGAGCTCTGAGTCTCCGCTCTCTCTCTGAGGATCGG 1280
Db 714 CTTGCGGAGGTCTGCGCACTGCGGCTCTCTGAGTCTCTAGCTGTTCTCTCAGGCGCTA 773
QY 1281 TCAACGAGCGGTCGCGCCCTCTGCA 1307
Db 774 GAGTCTCCGCCCCAGACAGCGGTTTCA 800

RESULT 6
ID ABK10801 standard; DNA; 2819 BP.
AC ABK10801;
XX
XX
21-MAY-2002 (first entry)
XX
XX Murine nucleosomal assembly protein IL2 (NapIL2 or Bpx) coding sequence.
XX
XX Murine nucleosomal assembly protein IL2; NapIL2; Bpx; nootropic;
KW

KW neurotropic; cytostatic; neural system defect; cancer;
KW neuronal cell tumour; gene therapy; neurodegenerative disease;
KW Parkinson's disease; Alzheimer's disease; spina bifida; anencephaly;
KW brain structure; brain organisation; X-chromosome linked disorder; rol;
KW nucleosome activity; cell cycle; cyclin; transcription factor control;
KW histone; gene; ds; mouse.
XX
XX Mus sp.
OS
XX
XX
PN WO200185995-A2.
XX
XX 15-NOV-2001.
XX
XX 04-MAY-2001; 2001WO-IB00960.
XX
XX 05-MAY-2000; 2000US-202111P.
XX
XX (INSP) INST PASTEUR.
XX (CNRS) CNRS CENT NAT RECH SCI.
XX
XX Avner P, Rogner UC, Spyropoulos D, Rougeulle C;
XX
XX WPI; 2002-075254/10.
XX
XX Screening neural system defects in mammal, comprises detecting a
XX modification of NapIL2 and/or NapIL2 gene chromosomal material, that
XX causes a loss of biological function and correlating the modification
XX with the defect -
XX
XX Disclosure; Fig 8; 83pp; English.

XX The invention describes a method of screening mammalian neural system
XX defects, comprising detecting a modification of the nucleosomal assembly
XX protein NapIL2 and/or NapIL2 gene in a human and/or mouse, where
XX modification is substitution, deletion, frame-shift, insertion aberrant
XX or altered epigenetic control, that causes a loss of function in the gene
XX and correlating the gene modification with a potential for a neural
XX system defect. The invention includes a neuronal cell containing a
XX recombinant chromosome on which NapIL2 is located. The cell is useful for
XX screening therapeutic compounds by introducing a compound to be screened
XX and correlating change in the proliferation of cells with the activity of
XX the compound, where change in proliferating cells is preferably a control
XX of cancer of neural cells. The NapIL2 promoter is useful for predicting
XX tumour of neural origin, and for genetic therapy of neurodegenerative
XX diseases such as Parkinson's, Alzheimer's diseases or accidents. NapIL2
XX is also implicated in spina bifida with or without anencephaly, loss of
XX brain structure/organisation, X-chromosome linked disorders and
XX inappropriate control of nucleosome activity in neurons, neuronal cell
XX cycle, cyclins, DNA binding transcription factors, histones and histone
XX shuttling. This sequence represents the murine nucleosomal assembly
XX protein IL2 (NapIL2 or Bpx) gene, described in the invention.
XX
SQ

Query Match 11.0%; Score 166.8; DB 24; Length 2819;
Best Local Similarity 64.9%; Pred. No. 1.7e-28;
Matches 322; Conservative 0; Mismatches 147; Indels 27; Gaps 4;
QY 827 CTTCTCTCATTTGTCGACATTTTGTAGTTACGTTTCTCTGAGCTCTCTGGAACCATAAAA 886
Db 5 CTTGCCCATTTTTTTCATTTCTCTGTCACCTGTTTCCATATAGCTCTTTTGAATTTATGAC 64
QY 887 GTATAATATCTGTTAAAAAGTTGGATGAATGAATGAACGCAATGGATTCCAGAAAA 946
Db 65 ATATAGTATCAGTTGAAAAACCGAATGAATGA-----TACTGATTTCTGCAAAA 113
QY 947 CTTCTGGGAGATGGCTAGAGGACGAGGAGGTGATGAATCAGCCATGTTTAGAGAG 1006
Db 114 TTCCACAGGCTATAGGTTGGAAGATGAGCCATAGTGGAGGAATCAGCCATATTAGAGAA 173
QY 1007 CCTGGGAAGGTGAGCAGAGTTGAAAACCTTGATAGATCTAATTAATTTACTGGCTCTGGGTT 1066
Db 174 TCTGGGAAGCAAGAGGTGTTGAAATTTTGAATTTTCACTACTAATTTACTGGCTCAGGATT 233

GenCore version 5.1.3
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OM nucleic - nucleic search, using sw model

Run on: January 8, 2003, 16:38:10 / Search time 3929 Seconds

(without alignments)
11258.912 Million cell updates/sec

Title: US-09-847-665-4

Perfect score: 1520
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Scoring table: IDENTITY_NUC

Gapop 10.0, Gapext 1.0

Searched: 2054640 seqs, 1451402878 residues

number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database:

GenBank1:
1: gb_ba:*
2: gb_hcg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
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35: em_hcg_rod:*
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37: em_hcg_vrt:*
38: em_sy:*
39: em_hcgo_hum:*
40: em_hcgo_mus:*
41: em_hcgo_other:*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1520	100.0	1520	6	AX327802
2	1520	100.0	3699	6	AX327804
3	1520	100.0	15378	9	AC004074
4	522.4	34.4	2720	6	AX327801
5	356	23.4	2515	9	AB027013
6	299.2	19.7	2524	9	BC026325
7	262	17.2	272	9	HS846R
8	233.8	15.4	1725	6	AX327799
9	233.8	15.4	210614	2	AL773526
10	233.8	15.4	214384	10	AJ421480
11	166.8	11.0	2819	6	AX327800
12	70.4	4.6	2226	3	MIDVTRN
13	69.8	4.6	154995	9	AC011979
14	69.6	4.6	152409	2	PFMAL1P1
15	69.4	4.6	108902	2	AC011430
16	68.6	4.5	123589	2	AC0104790
17	68.6	4.5	172758	2	AC022553
18	68.4	4.5	224448	2	PFMAL4P4
19	68.4	4.5	349980	6	AX344563
20	68.2	4.5	147192	2	AC116925
21	68.2	4.5	245802	2	AC006279
22	68	4.5	118642	2	AC126282
23	68	4.5	159712	2	AP000792
24	68	4.5	193227	2	AP000792
25	67.4	4.4	2426	8	SDU49822
26	67.4	4.4	349980	6	AX344555
27	67.2	4.4	169841	2	AC073888
28	67.2	4.4	177476	2	AC103159
29	67.2	4.4	199200	2	AL359633
30	67	4.4	213535	6	AX197417
31	67	4.4	213535	6	AX223856
32	66.6	4.4	97683	2	AC116548
33	66.4	4.4	143331	9	AC091214
34	66	4.3	86827	3	PFMAL3P5
35	65.4	4.3	10347	3	PFVALR
36	65.4	4.3	105686	9	AC012072
37	65.4	4.3	168698	9	AC068138
38	65.4	4.3	170796	2	AC069416
39	65.2	4.3	17967	6	AX345917
40	65.2	4.3	148202	2	AL772396
41	65.2	4.3	209856	2	AC117020
42	64.8	4.3	6419	6	AX345169
43	64.8	4.3	163964	2	AC106154
44	64.8	4.3	206059	2	AC127383
45	64.6	4.2	103344	9	HS1100E15

ALIGNMENTS

RESULT 1	AX327802	1520 bp	DNA	linear	PAT 07-JAN-2002
LOCUS	AX327802				
DEFINITION	Sequence 4 from Patent WO0185995.				
ACCESSION	AX327802				
VERSION	AX327802.1				
KEYWORDS	GI:18098075				
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;				
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.				
TITLE	Avner, P., Rogner, U.C., Spyropoulos, D. and Rougulle, C.				
	Identification of neural defects associated with the nucleosomal				
	assembly protein 112 gene				

Query Match	100.0%	Score 1520;	DB 6;	Length 3699;
Best Local Similarity	100.0%	Pred. No. 4.1e-312;		

Matches 1520; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Oy 1 ACTTAAAGAAAAATTTATCTATTAACGACAGATTTAGAAAATTAACAAATATGT 60
Db 1 ACTTAAAGAAAAATTTATCTATTAACGACAGATTTAGAAAATTAACAAATATGT 60
Oy 61 AAAAGTTTAAATCTGTGATAGTAACAAAATCTTTAAATCTGAAAATAATATGCT 120
Db 61 AAAAGTTTAAATCTGTGATAGTAACAAAATCTTTAAATCTGAAAATAATATGCT 120
Oy 121 TAAATTTTAAAAATTTGTCATTAATAATGATCCAAAGTTGAAATATGAAACAAATA 180
Db 121 TAAATTTTAAAAATTTGTCATTAATAATGATCCAAAGTTGAAATATGAAACAAATA 180
Oy 181 AACCTCAACCAATATTAATAGAGAGAAATTTTAATTAATCTCAAAAGCTTCCATCTA 240
Db 181 AACCTCAACCAATATTAATAGAGAGAAATTTTAATTAATCTCAAAAGCTTCCATCTA 240
Oy 241 TAAATCATTAATAATAGTTTAAACATTTCTTTAATGCTGAGATTTAGATTTTCCAA 300
Db 241 TAAATCATTAATAATAGTTTAAACATTTCTTTAATGCTGAGATTTAGATTTTCCAA 300
Oy 301 TTAATCAAAAGCATCAGCAAAATGTTATGATTTCTAAGATTAACATTAATCTTCAATT 360
Db 301 TTAATCAAAAGCATCAGCAAAATGTTATGATTTCTAAGATTAACATTAATCTTCAATT 360
Oy 361 TGGCTTTGATATATATATATTTCTAAGCGCTGTTAAAGCAGCATTAAGAGAGAG 420
Db 361 TGGCTTTGATATATATATATTTCTAAGCGCTGTTAAAGCAGCATTAAGAGAGAG 420
Oy 421 CAGAAAGCATATGAGGAGCTGGGTTATTTTAAGCCAGGCACTGGTTAATGTGTT 480
Db 421 CAGAAAGCATATGAGGAGCTGGGTTATTTTAAGCCAGGCACTGGTTAATGTGTT 480
Oy 481 AATGCTGATATGTTTACTAGTCACTGATGTTGATATACCAATACATTAATTTTATCA 540
Db 481 AATGCTGATATGTTTACTAGTCACTGATGTTGATATACCAATACATTAATTTTATCA 540
Oy 541 GGGCCCTATGCGCCCACTGCGATCGGACTTCCCTCCCTCCCTCCAGAGAAATGTT 600
Db 541 GGGCCCTATGCGCCCACTGCGATCGGACTTCCCTCCCTCCCTCCAGAGAAATGTT 600
Oy 601 CGAGAAATTTTCAACCTAAATCATATAGCTTGTAAGAAAATACGAAACATATATAG 660
Db 601 CGAGAAATTTTCAACCTAAATCATATAGCTTGTAAGAAAATACGAAACATATATAG 660
Oy 661 AATATTTAAATACGACGCGCACTTAAGACATCAATGCTAATTTCTGTGTTTAA 720
Db 661 AATATTTAAATACGACGCGCACTTAAGACATCAATGCTAATTTCTGTGTTTAA 720
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Db 721 ATCTTTAAGCGTTGTTTATCACTCTTCCACATCCCTCCCTCCCTCCAGAGTCC 780
Oy 781 CGATCTAAATTCAAAGAGATTTAGATGATGGTGGTGGTGGTGGTGGTGGTGGTGGT 840
Db 781 CGATCTAAATTCAAAGAGATTTAGATGATGGTGGTGGTGGTGGTGGTGGTGGTGGT 840
Oy 841 CGACATTTTATAGTTTCTCTGAGCTCTCTGAAAGCATTAAGTAAATATCTGTT 900
Db 841 CGACATTTTATAGTTTCTCTGAGCTCTCTGAAAGCATTAAGTAAATATCTGTT 900
Oy 901 AAAAGTTGATATGAATGAATGAATGAATGAATGAATGAATGAATGAATGAATGAAT 960
Db 901 AAAAGTTGATATGAATGAATGAATGAATGAATGAATGAATGAATGAATGAATGAAT 960
Oy 961 GGGCTAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1020
Db 961 GGGCTAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1020
Oy 1021 CAGAGTTGAAAACTGATGATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 1080
Db 1021 CAGAGTTGAAAACTGATGATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 1080

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Oy 1081 TTGCAAAATGAGATTAGACATAGTTGGAGAGGAGAGAGAGAGAGAGAGAGAGAGAG 1140
Db 1081 TTGCAAAATGAGATTAGACATAGTTGGAGAGGAGAGAGAGAGAGAGAGAGAGAGAG 1140
Oy 1141 TTGCACTAGAAATTTTAAAGCAAGTATAGTGGTATCATACTGCGGACCGTTTTT 1200
Db 1141 TTGCACTAGAAATTTTAAAGCAAGTATAGTGGTATCATACTGCGGACCGTTTTT 1200
Oy 1201 TCTTGACAGAGTACTCTCTGCGAGAGAGAGTCTGCCACTGACAGTCTCTGAGTCCG 1260
Db 1201 TCTTGACAGAGTACTCTCTGCGAGAGAGAGTCTGCCACTGACAGTCTCTGAGTCCG 1260
Oy 1261 GCTCTCTCTGACAGATCGGTCAACGAGCGGCGGCTCTGACCCAGCCAGGTC 1320
Db 1261 GCTCTCTCTGACAGATCGGTCAACGAGCGGCGGCTCTGACCCAGCCAGGTC 1320
Oy 1321 GCGCATGCTTCAAGTCGGTCTCTCAAAAGCTCAGACATCTTTATCCCGAGACCTG 1380
Db 1321 GCGCATGCTTCAAGTCGGTCTCTCAAAAGCTCAGACATCTTTATCCCGAGACCTG 1380
Oy 1381 GATGTCGTCCTCTGACGTCGAGCGGCGCATGCTAGGTCGACACCGCGCTTGATAT 1440
Db 1381 GATGTCGTCCTCTGACGTCGAGCGGCGCATGCTAGGTCGACACCGCGCTTGATAT 1440
Oy 1441 TTGGTGAGCTCTTCTCTGAGAGGTTTGCTCCGATCTCTGTAGGACACCTTAGG 1500
Db 1441 TTGGTGAGCTCTTCTCTGAGAGGTTTGCTCCGATCTCTGTAGGACACCTTAGG 1500
Oy 1501 CGGTACGGTCTTTGAAAA 1520
Db 1501 CGGTACGGTCTTTGAAAA 1520

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RESULT 3
AC004074 153578 bp DNA linear PRI 12-APR-1998
DEFINITION Homo Sapiens Chromosome X clone bXMD759, complete sequence.
AC004074
VERSION
KEYWORDS
SOURCE
ORGANISM
Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 153578)
Chen, R., Brownstein, B. H., States, D. J., Schlessinger, D. and
Mazzarella, R.
Mazzarella, R.
Direct Submission
Unpublished (1997)
2 (bases 1 to 153578)
Brownstein, B. H., States, D. J. and Mazzarella, R.
Submitted (29-JUN-1998) Center for Genetics in Medicine, Box 8232,
Washington University School of Medicine, 4566 Scott Avenue, St.
Louis, MO 63110, USA
3 (bases 1 to 153578)
Brownstein, B. H., States, D. J. and Mazzarella, R.
Submitted (12-APR-1998) Center for Genetics in Medicine, Box 8232,
Washington University School of Medicine, 4566 Scott Avenue, St.
Louis, MO 63110, USA
On Apr 13, 1998 this sequence version replaced gi:2822135.
Current status of this project is available at:
'http://www.ibc.wustl.edu/cgm/seq_projects.html'
Submitted by:
Elison Chen,
Advanced Center for Genetic Technology,
Applied Biosystems Division of Perlin Elmer Corp.,
850 Lincoln Center Drive,
Foster City, CA 94404 USA
e-mail: elison@genseq.apltdio.com

REFERENCE 1
 AUTHORS Ayner, P., Rogner, U.C., Spyropoulos, D. and Rougeulle, C.
 TITLE Identification of neutral defects associated with the nucleosomal
 JOURNAL assembly protein 112 gene
 Patent: WO 018595-A 3 15-NOV-2001.
 INSTITUT PASTEUR (FR) ; CENTRE NATIONAL DE LA RECHERCHE
 SCIENTIFIQUE (CNRS) (FR)
 FEATURES Location/Qualifiers
 source 1..2720
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 BASE COUNT 808 a 461 c 645 g 806 t
 ORIGIN

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 Best Local Similarity 99.8%; Pred. No. 1.7e-100;
 Matches 523; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 997 TGTAGAGAGCTGGAGAGTGGAGAGTGAAGAACTTGATGATCTAATAATTTACTG 1056
 1 TGTAGAGAGCTGGAGAGTGGAGAGTGAAGAACTTGATGATCTAATAATTTACTG 60
 QY 1057 GCTCTGGGTTTGTCACTCACTACATGAGCAAAATGAGATTAGACATGTTGGAGAG 1116
 DB 61 GCTCTGGGTTTGTCACTCACTACATGAGCAAAATGAGATTAGACATGTTGGAGAG 120
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 DB 121 GAAGAGAGTGAAGAGAGATCTAATTTGCACTTAAGAAATTTAGGCAAGTATAGCTGGCT 180
 QY 1177 AATGATAGTGGAGAGAGCTTTTCTTGACAGAGAGTGGTGGAGAGAGTCTGCC 1236
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 DB 241 CACTGAGCTCTCTGAGCTCTCCGCTCTCTCTGACAGATCGGTCAACGAGCGCTGCC 300
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 QY 1357 CATCTTTATCCCGAGAGAGCTGGATGCTTCCCTCAGTCCGAGAGCACTGCTAGG 1416
 DB 361 CATCTTTATCCCGAGAGAGCTGGATGCTTCCCTCAGTCCGAGAGCACTGCTAGG 420
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 421 TCCGACACCGCGCTTGTATTTGCGTGAATTTCTGTTGAGGTTTGTCTCC 480
 QY 1477 GATCTGTGTGAGAGCACTTAGCGGTGACGCTCTTGA AAA 1520
 DB 481 GATCTGTGTGAGAGCACTTAGCGGTGACGCTCTTGA AAA 524

RESULT 5
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 DEFINITION Homo sapiens mRNA for Nucleosome Assembly protein 1-like 2,
 complete cds.
 ACCESSION AB027013
 VERSION AB027013.1 GI:5931609
 KEYWORDS Nucleosome Assembly Protein 1-like 2.
 SOURCE Homo sapiens
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
 Sasaki, N., Hattori, A., Hayashi, A., Kozuma, S., Muramatsu, M.,
 Miyajima, N. and Saito, T.
 Nucleosome Assembly Protein 1-like 2 (Brain-Specific Protein,
 X-linked)
 JOURNAL Published Only in Database (1999)

REFERENCE 2 (bases 1 to 2515)
 AUTHORS Seki, N., Hattori, A., Hayashi, A., Kozuma, S., Muramatsu, M.,
 Miyajima, N. and Saito, T.
 TITLE Direct Submission
 JOURNAL Submitted (07-MAY-1999) Toshiyuki Saito, National Institute of
 Radiological Sciences, Genome Research Group, Anagawa 4-9-1,
 Inage-Ku, Chiba 263-8555, Japan (E-mail: t_saito@nirs.go.jp,
 Tel:81-43-206-3135, Fax:81-43-251-9818)
 FEATURES Location/Qualifiers
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 /db_xref="taxon:9606"
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 BASE COUNT 742 a 432 c 589 g 752 t
 ORIGIN

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 Best Local Similarity 100.0%; Pred. No. 3.4e-65;
 Matches 356; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1165 TGATAGCTGGAATCATCTGCGACCGTTTCTTGAGAGAGTACGCTTGGG 1224
 DB 1 TGATAGCTGGAATCATCTGCGACCGTTTCTTGAGAGAGTACGCTTGGG 60
 QY 1225 AGAGAGCTGGCCAGTCACTGAGCTCTGAGTCTCCGCTCTCTGAGAGATCGGTCA 1284
 DB 61 AGAGAGCTGGCCAGTCACTGAGCTCTGAGTCTCCGCTCTCTGAGAGATCGGTCA 120
 QY 1285 CGCAGCGTGGCGGCTCTGACAGCCAGAGTGGCACTGCTTCAAGTCCGTTCTCA 1344
 DB 121 CGCAGCGTGGCGGCTCTGACAGCCAGAGTGGCACTGCTTCAAGTCCGTTCTCA 180
 QY 1345 AAGCTCAGACCATCTTTATCCCGAGAGAGCTGATCGTCCCTCAGTCCGAG 1404
 DB 181 AAGCTCAGACCATCTTTATCCCGAGAGAGCTGATCGTCCCTCAGTCCGAG 240
 QY 1405 GCCACTGCTAGAGTCCGACACCGCGCTTGTGATATTTCCGTGAGTCTTTCTGTGAG 1464
 DB 241 GCCACTGCTAGAGTCCGACACCGCGCTTGTGATATTTCCGTGAGTCTTTCTGTGAG 300
 QY 1465 GTTGTGCTCCGATCTCTGTGAGTACACCTTAGCGGTGAGGCTCTTGA AAA 1520
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RESULT 6
 LOCUS BC026325 2524 bp mRNA linear PRI 08-APR-2002
 DEFINITION Homo sapiens, nucleosome assembly protein 1-like 2, clone MGC:26243
 IMAGE:4798964, mRNA, complete cds.
 ACCESSION BC026325
 VERSION BC026325.1 GI:20070943
 KEYWORDS MGC.
 SOURCE Homo sapiens
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
 Strausberg, R.

Thu Jan 9 09:34:26 2003

TITLE
JOURNAL

REMARK
COMMENT

Direct Submission
Submitted (02-APR-2002) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
NIH-MGC Project URL: <http://mhc.nci.nih.gov>
Contact: MGC help desk
Email: cgabbs@mail.nih.gov
Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
cDNA Library Preparation: Michael J. Brownstein (NHGRI) & Shiraki
Toshiyuki and Piero Carninci (RIKEN)
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Sequencing Group at the Stanford Human Genome
Center, Stanford University School of Medicine, Stanford, CA 94305
Web site: <http://www-shgc.stanford.edu>
Contact: (Dickson, Mark) mcd@paxil.stanford.edu
Dickson, M., Schmutz, J., Grimwood, J., Rodriguez, A., and Myers,
R. M.

Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Series: IRAC Plate: 32 Row: g Column: 23
This clone was selected for full length sequencing because it
passed the following selection criteria: matched mRNA gi: 11415047.

FEATURES
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CDS

BASE COUNT 790 a 420 c 577 g 737 t

ORIGIN

Query Match 19.7%; Score 299.2; DB 9; Length 2524;
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Matches 30; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1217 GCTTGGCGAGGAGTCTGCCCATCGAGTCTCTCGAGTCTCGGCTCTCTCTCGAGG 1276
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QY 1277 TCGGTCAACGCGCGTCCGCGCCTCTGCACCCAGCCAGGTCGCCACTGCTTCAGTCC 1336
Db 61 TCGGTCAACGCGCGTCCGCGCCTCTGCACCCAGCCAGGTCGCCACTGCTTCAGTCC 120

QY 1337 GGTTCCTCAAGCCCTCAGCACCATCTTTATTCGCCGAGCAGCTGGATCGCTTCCTCA 1396
Db 121 GGTTCCTCAAGCCCTCAGCACCATCTTTATTCGCCGAGCAGCTGGATCGCTTCCTCA 180

QY 1397 GTCCGAGCGCAGCTGAGTTCGACACCGCGCTTCTGATATTTTCGGTGAAGTCTTTTC 1456
Db 181 GTCCGAGCGCAGCTGAGTTCGACACCGCGCTTCTGATATTTTCGGTGAAGTCTTTTC 240

QY 1457 CTGTGGAGGTTTGGTCTCCCGATCTCTGTGGTAGCCACCTTAGCGGTGACGTCCTTTC 1516

Db 241 CTGTGGAGGTTTGGTCTCCCGATCCCTGTGGTAGCCACCTTAGCGGTGACGTCCTTTC 300

QY 1517 AAAA 1520
|||||

Db 301 AAAA 304

RESULT 7

HS84H6R
LOCUS
DEFINITION
H. sapiens CpG island DNA genomic MseI fragment, clone 84h6, reverse
read cp984h6.rcl1a.
263470
VERSION
263470.1 GI:1035848
KEYWORDS
CpG island; genomic MseI fragment.
SOURCE
Homo sapiens
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE
AUTHORS
TITLE
JOURNAL
DIRECT SUBMISSION
Submitted (16-OCT-1995) The Sanger Centre, Hinxton, Cambridgeshire,
CB10 1RQ, England. E-mail contact: humquery@sanger.ac.uk
2 (bases 1 to 272)

REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
COMMENT
Cross, S.H., Charlton, J.A., Nan, X. and Bird, A.P.
Purification of CpG islands using a methylated DNA binding column
Nat. Genet. 6 (3), 236-244 (1994)
8012384

Vector: pGEM-52f (-)
Clones are available from the UK MRC Human Genome Mapping Project
Resource Centre, Hinxton, Cambridgeshire CB10 1RQ, UK. See URL:
<http://www.hgmp.mrc.ac.uk/> for details
or contact: biohelp@hgmp.mrc.ac.uk.

FEATURES
source

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Best Local Similarity 96.3%; Pred. No. 2.5e-45;
Matches 26; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 902 AAAGTTGGATGAATGAACCAATGAACGCAATGGATTCCAGAAACTCTGCGGAGATGG 961
Db 1 AAAGTTGGATGAATGAACCAATGAACGCAATGGATTCCAGAAACTCTGCGGAGATGG 60

QY 962 GCTAGAGGACGAGAGGAGTGGATGAATCAGCCATGTTAGAGAGCTCGGAAGGTGAGC 1021
Db 61 GCTAGAGGACGAGAGGAGTGGATGAATCAGCCATGTTAGAGAGCTCGGAAGGTGAGC 120

QY 1022 AGAGTTGAAACTTGTAGATCTAATAATTTACTGGCTCTGGGTTTTCAGTCACTACAT 1081
Db 121 AGAGTTGAAACTTGTAGATCTAATAATTTACTGGCTCTGGGTTTTCAGTCACTACAT 180

QY 1082 TGCAGCAAAATGAGATTAGAGCATAGTTGTGGAGGAGGAGGTGAGCGCAATCTATT 1141
Db 181 TGCAGCAAAATGAGATTAGAGCATAGTTGTGGAGGAGGAGGTGAGCGCAATCTATT 240

QY 1142 TGCACCTAGTAATTTTAGGCAAGTGATAGCTG 1173
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RESULT 8

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DEFINITION   Sequence 1 from Patent WO0185995.
ACCESSION    AX227799
VERSION      AX227799.1  GI:18098072
KEYWORDS
SOURCE       Mus sp.
ORGANISM     Mus sp.
             Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
             Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE
AUTHORS      1
TITLE        Aymer, P., Rogner, U.C., Spyropoulos, D. and Rougeulle, C.
             Identification of neutral defects associated with the nucleosomal
             assembly protein 112 gene
JOURNAL      Patent: WO 0185995-A 1 15-NOV-2001;
             INSTITUT PASTEUR (FR) ; CENTRE NATIONAL DE LA RECHERCHE
             SCIENTIFIQUE (CNRS) (FR)
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             /db_xref="taxon:10095"
BASE COUNT   428 a 443 c 391 g 463 t
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Query Match 15.4%; Score 233.8; DB 6; Length 1725;
Best Local Similarity 65.1%; Pred. No. 2.8e-39;
Matches 486; Conservative 0; Mismatches 222; Indels 39; Gaps 8;

QY 580 CTCCTCCACAGAAATGTTTCAGAAATTTTCAACCTTAATCATATAGCTTGAA 639
DB 74 CTCCTCATAGAAATATATAGAAATTTTCCATTAAATCACAGGTTGGAAA 133
QY 640 ATACCGAACAACTATATAGAAATTTTAAATTAATCTGAC---ACGCCACTTAAGACA 695
DB 134 TTAGAGA-AACAGAGGTACAGAAATTTAAACACTGTGATCATATCAAAAGCCA 192
QY 696 TCAAGCTTAATCTGTGTTTATCTTTAGAGCGTTTGTATCAGCTCTTCCACA 755
DB 193 CTTAGCTTAATTTTGTATTTTAACTCAAGATCTCTTGTGGCTCTCCACTA 252
QY 756 TCACCTCTCCCTCCCAAGCGTCCCGATTAATCAAGAGATTTAGAGATGGGT 815
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DB 311 CAGTACCTGCCATTTTTCATTG---CTGTGACGTGTTCCATATGCTCTTTG 365
QY 876 AAAGCATAAAGTAAATCTGTTAAAGTTGATGATGAACTAAATGAACCAATGG 935
DB 366 AATATATGAACATATAGATCAGTTGAACGGAATGATGA-----TACTGCA 414
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DB 415 TTTCGCAAAATTCACAGGCTATAGGTGAGAGATAGCATAGGTGAGGAATACGCC 474
QY 996 ATGTTAGAGACCTCGGAGAGTGAAGAGTTGAAAATTGATAGATCTTAATTTACT 1055
DB 475 ATATTAAGATCTGGAAGAGCAAGAGGTGTAATTTGATTCTACTAATTTACT 534
QY 1056 GGGCTGGGTTTGTCACTCACTACATGACGAATGAGTTGACATAGTTGGAG 1115
DB 535 GGGCTCAGAGTTTGTCACTCACTGAGCTGCAAAATGAGATTGAGAAAGCTCGGAG 594
QY 1116 GGAAGAGGTGAGGAGCAATCTATTGCACT-AGAAATTTTAGCAAGTATAGTGC 1174
DB 595 GGAAGG-GGTGAGGAGCAACCTGACATACCTTAATAAAGAGTGAGACAACTGC 653
QY 1175 GTAATCACTGCGGACCGTTTTTTCTT-----GCAGCATGAGTGGCTT 1220
DB 654 GTAATCACTGCGGACCGTTTTTTCTT-----GCAGCATGAGTGGCTG 713
QY 1221 GCGAGAGAGTCTGCCACTGACGCTCTCTGAGATCTCCGCTCTCTCTGAGGATGG 1280

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DB 714 CTGGGAGAGTGTGCCCTGAGCTGCTCTGCACTCTGCTGTTCTTCAAGGGCTTA 773
QY 1281 TCAACGACCGCTGCGCGCCCTCTGCA 1307
DB 774 GAGTCTCGCCACAGACCGCGTTTCA 800

RESULT 9
LOCUS      AL773526/c
DEFINITION Mus musculus chromosome X clone RP23-181M13, *** SEQUENCING IN
ACCESSION  AL773526
VERSION    AL773526.5  GI:22204627
KEYWORDS  HTGS_PHASE1; HTGS_ACTIVEFIN; HTGS_DRAFT; HTGS_FULLTOP.
SOURCE     house mouse.
ORGANISM  Mus musculus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
AUTHORS    Direct Submission
TITLE      Submitted (16-AUG-2002) Wellcome Trust Sanger Institute, Hinxton,
           Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
           humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
           On Aug 11, 2002 this sequence version replaced gi:121621733.
JOURNAL    ----- Genome Center
           Center: Wellcome Trust Sanger Institute
           Center code: SC
           Web site: http://www.sanger.ac.uk
           Contact: humquery@sanger.ac.uk
           Project Information
           Center project name: BM181M13
           ----- Summary Statistics
           Assembly program: XGAP4; version 4.5
           Chemistry: Dye-terminator; 100% of reads
           Consensus quality: 209617 bases at least Q40
           Consensus quality: 209945 bases at least Q30
           Insert size: 210214; sum-of-contigs
           Insert size: 194526; 6.8% error; agarose-fp
           Quality coverage: 9.25x in Q20 bases; sum-of-contigs Quality
           coverage: 10.05x in Q20 bases; agarose-fp

* NOTE: This is a 'working draft' sequence. It currently
* consists of 5 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 23338: contig of 23338 bp in length
* 23339 23438: gap of 100 bp
* 23439 99523: contig of 76085 bp in length
* 99524 99623: gap of 100 bp
* 99624 144306: contig of 44683 bp in length
* 144307 144406: gap of 100 bp
* 144407 201734: contig of 57328 bp in length
* 201735 201834: gap of 100 bp
* 201835 210614: contig of 8780 bp in length.
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Thu Jan 9 09:34:26 2003

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Best Local Similarity 65.1%; Pred. No. 3.8e-39;
Matches 486; Conservative 0; Mismatches 222; Indels 39; Gaps 8;
QY 590 CTCCTCCACAGGAATGTTTCGAGATTTTCAACCTAAATCATATAGCTTGTGAAA 639
Db 52001 CTTCTCATCAGGAATATTATGAGATTTTCCCAATTAAATCACACAGGTTGTGAAA 51942
QY 640 ATACCGACAAACATAATAGATATTAAATACTGAC-----ACGCCACTAAAGACCA 695
Db 51941 TTACAGA-AACAGGGTACAGATATTAAACCAGTGTACATCATCAAGGCCA 51883
QY 696 TCAGTGTAAATCTCGTGTGTTTAACTTTGAGCGTTTGTATCAGCTCTTCCACCA 755
Db 51882 CCTATGCTTATTTTGGTAATTTTAAACCTCAAGGATCTTTTGTGGGCTCTCCACTA 51823
QY 756 TCACCTCTCCCTCCAGGTCCTCCGATCTTAAATCAAGAGATTAATTTAGATGGGT 815
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Db 51764 CAGTACCTTGCCCAATTTTCTTCAATTC-----CTTGTCACTGTTTCCATATAGCTTTTG 51710
QY 876 AAGCATAAAGTATATATATCTGTTAAAGTTGATGATGAATGAATGAATGAATGAATGA 935
Db 51709 AAATTATGACATATATAGTATCAGTTGAAACCGAATGAATGA-----TACTGCA 51661
QY 936 ATTCCAGAAATCTCTCGGAGATGGCTTAGAGCAGGAGGAGGTGATGAATCAGCC 995
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QY 1175 GTAATCATCTGCGGACCGTTTCTT-----GCGAGTATGCTGCTT 1220
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Db 51361 CTTGCGGAGGTGTCGCCACTGCGGCTCTCTGACGCTCTCTGCTCTCTCTCTCTCTCTCTCT 51302
QY 1281 TCAAGCAGCGCTGCGGCCCTCTGCA 1307
Db 51301 GAGTCTCCGCCACGACAGCGCGTTTCA 51275
RESULT 10
AJ421480 214384 bp DNA linear ROD 12-JUN-2002
LOCUS
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Best Local Similarity 65.1%; Pred. No. 3,8e-39;
Matches 486; Conservative 0; Mismatches 222; Indels 39; Gaps 8;

QY 580 CTCCTCCAGCAATATGTTCCAGATTTTTCACCTAAATCATATAGCTTGCAAA 639
DB 161158 CTTCTCATCGAGAAATATATGAGATTTTCCATTAAATCACAGGTGTGAAA 161217
QY 640 ATACGCAAAACATATATGAAATATTAATACTGAC---ACCCACCTTAACACA 695
DB 161218 TTACAGA-AACAGAGGATGAGATATTTAAACACCTGTCATCATCAAAAGCCA 161276
QY 696 TCACTGTATTCCTGATGTTTATTCCTTGAAGGTTTGTATTCACCTTCACCA 755
DB 161277 CTTATGCTATTTTGTATTTTAAACCTCAAGGATCTTTGGGGCTCTCCACA 161336
QY 756 TCCACCTCTCCCTCCCGAGTCCGATCTAAATCAAGAGATTGATTGATGGGT 815
DB 161337 CCTCTCTCT--CTTCCAGAGCTCAGGTTATTAACCAAGGATGACTAAAGCAATC 161394
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QY	816	GGGTGCTTGTCTTCTCATTCTTCGACATTTTAGTTACGTTTCTCTGAGCTCTCTGG	875
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QY	936	ATTCCAGAAAACCTCTGGGAGATGGCTAGAGGACGAGGAGGTGGATGAATCAGCC	995
Db	161499	TTTCTGCAAAATTCACAGGCTATAGGGTGAAGATGAGCCATAGGTGGAGGAATCAGCC	161558
QY	996	ATGTTAGAGAGCTGGGAAGGTGAGCAGAGTTGAAAACCTTGATAGATCTTAATATTACT	1055
Db	161559	ATATTAGAGATCTGGGAAGGACAGAGGTGTTGAAATTTTGAATTTCTACTAATATTACT	161618
QY	1056	GGCTCTGGGTTTGTCACTCACTACATTGCAGCAATCAGATTAGAGCATAGTTGTGGAG	1115
Db	161619	GGCTCAGGATTTGTCAATCACTGCAGCTGGCAATCAGATTAGAGAGATCTCTGGAG	161678
QY	1116	GGAGGAGTGCAGCAGCAATCTATTGTCACCT-AGAAATTTTAGGCAAGTGTAGCTGC	1174
Db	161679	GGAGG-GGTGACGAGCAACTGCATACACTTAAAAAAAAGAGCTGAGAGCAACTGC	161737
QY	1175	GTAATCATATCTGGGACCGTTTTTTTCTT-----GCAGCAGTAGCTGCTT	1220
Db	161738	GTAATCATATCTGGGACCGATTCTCCATCCCTCCGCCCGGAGTGTGTCGAGCACTG	161797
QY	1221	GGGAGGAGTCTGCCACTGCAGCTCTGCAGTCTCGGCTCTCTCTCTCTCGCAGGATCGG	1280
Db	161798	CTTGGGAGGTCTGCCACTGCAGCTCTCGGCTCTCTGCAGTCTCTAGCTGTTCTCTTCAGGCGCTA	161857
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Db	161858	GAGTCTCCGCCGACAGCGCGTTTCA	161884
RESULT 11			
LOCUS	AX327800	2819 bp	DNA linear PAT 07-JAN-2002
DEFINITION	Sequence 2 from Patent WO0185995.		
ACCESSION	AX327800		
VERSION	AX327800.1	GI:18098073	
SOURCE	Mus sp.		
ORGANISM	Mus sp.		
REFERENCE	Ayner, P., Rognier, U.C., Spyropoulos, D. and Rougeulle, C.		
AUTHORS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.		
TITLE	Identification of neural defects associated with the nucleosomal assembly protein 12 gene		
JOURNAL	Patent: WO 0185995-A 2 15-NOV-2001; INSTITUT PASTEUR (FR) ; CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE (CNRS) (FR)		
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Best Local Similarity	64.9%;	Pred. No. 4.6e-25;	
Matches 322;	Conservative	0; Mismatches 147;	Indels 27; Gaps 4;
QY	827	CTTCTCTCATTTGTCGACATTTTAGTTAGCTTTTCTCTGAGCTCTCTGGAAGCATAAAA	886
Db	5	CTTGGCCATTTTTCATTCTCTGTCACGTGTTCCATATAGCTCTTTGAAATTTATGAC	64
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RESULT 2
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LOCUS 367 bp mRNA linear EST-20-Apr-1997
DEFINITION EST27025 Cerebellum II Homo sapiens cDNA 5' end, mRNA sequence.
VERSION AA324132.1 GI:1976479
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 367)
AUTHORS Adams, M.D., Karlavage, A.R., Fleischmann, R.D., Fuldner, R.A., Bult
C.J., Lee, N.H., Kirkness, E.F., Weissbrock, K.G., Gocayne, J.D., White
O., Sutton, G., Blake, J.A., Brandon, R.C., Man-Wai, C., Clayton, R.A.,
Cline, T.R., Cotton, M.D., Barle-Hughes, J., Fine, L.D., Fitzgerald,
L.M., Fitzhugh, W.M., Fritchman, J.L., Geoghegan, N.S., Glodok, A.,
Gnehm, C.L., Hanna, M.C., Hedblom, E., Hinkle, P.S. Jr., Kelley, J.M.,
Kelley, J.C., Liu, L.-I., Marmaros, S.M., Merrick, J.M.,

TITLE
JOURNAL
MEDLINE
COMMENT

Moreno-Palanges, R.F., McDonald, L.A., Nguyen, D.T., Pelligri,
Phillips, C.A., Ryder, S.B., Scott, J.L., Saudex, D.M., Shirley, R.,
Small, K.V., Spriggs, T.A., Uteerback, T.R., Weidman, J.P., Li, Y.,
Bedarick, D.P., Cao, L., Cepeda, M.A., Coleman, T.A., Collins, E.J.,
Dime, D., Feng, D.-F., Ferris, A., Fischer, C., Hastings, G.A., H.
Hu, J.S., Greene, J.M., Gruber, J., Hudson, P., Kim, A.K., Kozak,
Kunach, C., Hungjun, J., Li, H., Meisner, P.S., Olsen, H., Raymont,
Wei, Y.F., Wing, J., Xu, C., Yu, G.L., Ruben, S.M., Dillon, P.J.,
Venter, J.C., Rosen, C.A., Haseltine, W.A., Fields, C., Fraser, C.M. and
Initial assessment of human gene diversity and expression pati
based upon 83 million nucleotides of cDNA sequence
Nature 377 (6547 Suppl.) 3-174 (1995)
96026280
Contact: Kerlavage, AR
Bioinformatics
The Institute for Genomic Research
9712 Medical Center Drive, Rockville, MD 20850 USA
Tel: 3018699056
Fax: 3018699423
Email: arkerlav@igrr.org
For clone availability, additional sequence and expression
information related to this EST, please check the TIGR Human (c
Index (<http://www.tigr.org/cdb/hgi/hgi.html>)
Seq primer: M13 Reverse.

FEATURES

source

1. 367
Location/Qualifiers

/organism="Homo sapiens"
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/clone_1ib="Cerebellum II"
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/dev_stage="adult"
/note="Organ: brain; Vector: pBluescript SK-; Site_1
EcorI; Site_2: XhoI"

BASE COUNT 54 a 113 c 92 g 98 t 10 others
ORIGIN

Query Match

Best Local Similarity 96.9%; Score 345.4; DB 9; Length 367;
Matches 346; Conservative 0; Mismatches 11; Indels 0; Gaps

QY 1164 GTGATAGCTGCGTATCATATGCGGACCGTTTCTTGGACAGTACTGCTGG 12;
DB 1 GTGATAGCTGCGTATCATATGCGGACCGTTTCTTGGACAGTACTGCTGG 60
QY 1224 GAGAGGTGCGGACCTGCACTCTCTGAGTCTCGGCTCTCTCTGAGAGTCTCA 12;
DB 61 GAGAGGTGCGGACCTGCACTCTCTGAGTCTCGGCTCTCTCTGAGAGTCTCA 121
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DB 121 AGCGACCGCTGCGCGCCCTCTGACCGGACGAGTGGCCACTGCTTCAAGTCTGCTC 186
QY 1344 AAAGCCTGAGCACTCTTTTATCCCGAGAGAGCTGAGTCTGCTTCCCTGAGTCTCGGA 14;
DB 181 AAAGCCTGAGCACTCTTTTATCCCGAGAGAGCTGAGTCTGCTTCCCTGAGTCTCGGA 24;
QY 1404 CGCCACTGCTGAGTCTGAGCAACGCGGCTCTGATTTTGGGAGTCTTTTCTGAGGA 146
DB 241 CGCCACTGCTGAGTCTGAGCAACGCGGCTCTGATTTTGGGAGTCTTTTCTGAGGA 306
QY 1464 GGTTCGCTCCCATCTCTGAGTACCACTTGAAGCGGTGAGGCTTTGAAA 1520
DB 301 GGTTCGCTCCCATCTCTGAGTACCACTTGAAGCGGTGAGGCTTTGAAA 357

RESULT 3

AA120587
LOCUS



[illegible]

COMMENT
On Apr 13, 1998 this sequence version replaced gi:282213
Current status of this project is available at:
'http://www.ibc.wustl.edu/cgm/seq_projects.html'
Submitted by:
Elison Chen,
Advanced Center for Genetic Technology,
Applied BioSystems Division of Perlin Biomer Corp.,
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Foster City CA 94404 USA
e-mail: elison@genseq.apltdio.com

and

Buddy Brownstein,
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Washington University School of Medicine, Box 8232
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and

David J. States,
Institute for Biomedical Computing
Washington University in St. Louis
700 South Euclid Ave.
St. Louis, MO 63108 USA
e-mail: states@ibc.wustl.edu.

FEATURES

source

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/db_xref="taxon:9606"
/chromosome="x"
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BASE COUNT 45069 a 28281 c 29096 g 50132 t
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Query Match 100.0%; Score 1520; DB 9; Length 153578;
Best Local Similarity 100.0%; Pred. No. 5.4e-312;

Matches 1520; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 ACTTAAAGAAAATTTTCTATTAACCTGACAGAAATTTAGAAATTAATACAAATATGT 60
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QY 61 AAACGTTTAAATCTGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 120
DB 136411 AAACGTTTAAATCTGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 136470
QY 121 TAAATTTTAAATAATGTTTCAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 180
DB 136471 TAAATTTTAAATAATGTTTCAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 136530
QY 181 AACCTGACCAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 240
DB 136531 AACCTGACCAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAATTAAT 136590
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DB 136831 AATTGCTGGATGTTTATAGTACAGTATGTTATACACCATATGATTTTTCATCACA 136890
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QY 601 CGAGAAATTTTCAACCTAAATCATATAGCTTGTGAAAAATACGACAAACATATATATAG 660
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DB 137131 CGATCTAAATCAAGAGATGATTTATAGATGAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGT 137190
QY 841 CGACATTTTATAGTATGTTTCTCTGAGCTCTCTGAGCTCTCTGAGCTCTCTGAGCTCTCTGAG 900
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DB 137251 AAAAGTTGATGATGAACTAATGAAAGCAATGGGATTCAGAAAACCTCTGCGGAGATG 137310
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DB 137311 GGCTAGAGAGCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 137370
QY 1021 CAGAGTTGAAAATTTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1080
DB 137371 CAGAGTTGAAAATTTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 137430
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DB 137431 TTGACGAAAATGAGATTGACATATGTTTGTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAT 137490
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DB 137551 TCTTGACAGCATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 137610
QY 1261 GCTCTCTCTCAGAGATCGGATCGACAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAG 1320
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DB 137671 GCACTGCTTCAAGTCCGATCTCAAAAGCTCAGACCAATCTTTATCCCGAGCAGCTG 137730
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Db 1501 CGTGTACGCTCTTTGAAAA 1520

RESULT 3

AC004074

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

HTG.

HTG.

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COMMENT

AC004074 153578 bp DNA linear PRI 12-APR-1998
Homo Sapiens Chromosome X clone bMXD759, complete sequence.
AC004074
AC004074.1 GI:3046270
HTG.
Homo sapiens.
Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 153578)
Chen, E., Brownstein, B.H., States, D.J., Schlessinger, D. and
Mazzei, R.
Direct Submission
2 (bases 1 to 153578)
Brownstein, B.H., States, D.J. and Mazzei, R.
Direct Submission
Submitted (29-JAN-1998) Center for Genetics in Medicine, Box 8232,
Washington University School of Medicine, 4566 Scott Avenue, St.
Louis, MO 63110, USA
3 (bases 1 to 153578)
Brownstein, B.H., States, D.J. and Mazzei, R.
Direct Submission
Submitted (12-APR-1998) Center for Genetics in Medicine, Box 8232,
Washington University School of Medicine, 4566 Scott Avenue, St.
Louis, MO 63110, USA
On Apr 13, 1998 this sequence version replaced gi:2822135.
Current status of this project is available at:
'http://www.ihc.wustl.edu/cgm/seq_projects.html'
Submitted by:
Elison Chen,
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and

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and

David J. States,
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Washington University in St. Louis
700 South Euclid Ave.
St. Louis, MO 63108 USA
e-mail: states@bc.wustl.edu
Location: Qnailifiers

FEATURES

source

1. 153578
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="X"
/clone="BMX0759"

BASE COUNT 45069 a 29281 c 29096 g 50132 t

ORIGIN

Query Match 100.0%; Score 1520; DB 9; Length 153578;
Best Local Similarity 100.0%; Pred. No. 5.4e-312;

Matches 1520; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ACTTAAGGAAAAATTTATCTATTAACCTGACGAAATTTAGAAATTAATACAAATATGT 60
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Db 136591 TAAATACATTAATCAATAGTTTAACCAATTTCTTAAATGCTGAGATTTAGATTTTCCAA 136591
QY 301 TTAATCTAAAAAGCATCAAGCAAAATTTATGATTTCTAAGAAATTAACATTAATTTCCATT 360
Db 136651 TTAATCTAAAAAGCATCAAGCAAAATTTATGATTTCTAAGAAATTAACATTAATTTCCATT 136651
QY 361 TGGCTTTGTAATATATATATATTTCTAAGCGCTGTTAAAGCAGCATTAAAGAGAGAG 420
Db 136711 TGGCTTTGTAATATATATATATTTCTAAGCGCTGTTAAAGCAGCATTAAAGAGAGAG 136711
QY 421 CAGAAGGAGATTTGGAGATGGGGTATTTATTAAGCAGGCAATGGTTAATTTGGT 480
Db 136771 CAGAAGGAGATTTGGAGATGGGGTATTTATTAAGCAGGCAATGGTTAATTTGGT 136771
QY 481 AATTGCTGTATGTTTACTAGTACAGTATGTAATCAACATATAGTTTTCATCA 540
Db 136831 AATTGCTGTATGTTTACTAGTACAGTATGTAATCAACATATAGTTTTCATCA 136831
QY 541 GGCCCTCATTCGCCCCACAGCCCATCGGACTTCCTCTCTCCCTCAACAGAAATGTTT 600
Db 136891 GGCCCTCATTCGCCCCACAGCCCATCGGACTTCCTCTCTCCCTCAACAGAAATGTTT 136891
QY 601 CGAAGATTTTCAACCTAAATCATATAGCTGTGAAAAATACGCAAACTAATATAG 660
Db 136951 CGAAGATTTTCAACCTAAATCATATAGCTGTGAAAAATACGCAAACTAATATAG 136951

QY 661 AATTTTAAATTAATCTGACAGCCCACTAAAGACATCAATGCTAAATTCCTGTTT 720
Db 137011 AATTTTAAATTAATCTGACAGCCCACTAAAGACATCAATGCTAAATTCCTGTTT 137011
QY 721 ATCTTTAAGCGTTTGTATATAGCTCTTCCACATCCACTCTCCCTCCAGAGTCC 780
Db 137071 ATCTTTAAGCGTTTGTATATAGCTCTTCCACATCCACTCTCCCTCCAGAGTCC 137071
QY 781 CGATCTAAATTAATCAAGATTAATTAAGATGGGAGGCTCTCTCTCAATGTT 840
Db 137131 CGATCTAAATTAATCAAGATTAATTAAGATGGGAGGCTCTCTCTCAATGTT 137131
QY 841 CGACATTTAGTAACTTTTCTCTGAGCTCTCTGAGAAAGATTAATATATCTGTT 900
Db 137191 CGACATTTAGTAACTTTTCTCTGAGCTCTCTGAGAAAGATTAATATATCTGTT 137191
QY 901 AAAAGTTGATGATGAATGAATGAATGAATGAATGAATGAATGAATGAATGAATGA 960
Db 137251 AAAAGTTGATGATGAATGAATGAATGAATGAATGAATGAATGAATGAATGAATGA 137251
QY 961 GGCTAGAGACAG 1020
Db 137311 GGCTAGAGACAG 137311
QY 1021 CAGAGTTGAAATCTGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1080
Db 137371 CAGAGTTGAAATCTGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 137371
QY 1081 TTGACGAAATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1140
Db 137431 TTGACGAAATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 137431
QY 1141 TTGACCTAGAAATTTTAAAGGAAATGATGATGATGATGATGATGATGATGATGAT 1200
Db 137491 TTGACCTAGAAATTTTAAAGGAAATGATGATGATGATGATGATGATGATGATGAT 137491
QY 1201 TTTGACAGAGTACCTGCTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1260
Db 137551 TTTGACAGAGTACCTGCTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 137551
QY 1261 GCTCTCTGACAGAGTACCTGCTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1320
Db 137611 GCTCTCTGACAGAGTACCTGCTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 137611
QY 1321 GCACTGCTTCAAGTCCGCTTCTCAAAAGCTCAGACCATCTTTATCCCGAGAGCTTG 1380
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QY 1381 GATGCTGTTCCCTCAAGTCCGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1440
Db 137731 GATGCTGTTCCCTCAAGTCCGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 137731
QY 1441 TTGCGTATGCTTTTCTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1500
Db 137791 TTGCGTATGCTTTTCTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 137791
QY 1501 CGGTATAGGCTCTTTGAAA 1520
Db 137851 CGGTATAGGCTCTTTGAAA 137851

RESULT 4
AX327801

